

SPECIFICATION

TITLE OF THE INVENTION

NON-MUCIN TYPE SYNTHETIC COMPOUNDS OR IT'S CARRIER CONJUGATED COMPOUNDS

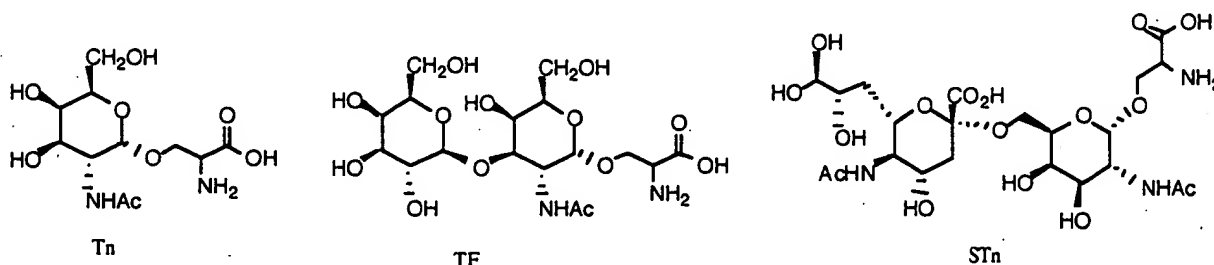
BACKGROUND OF THE INVENTION

1.The Subject of the Invention

The present invention concerns non-mucin type synthetic compounds which were linked to carrier, that is, non-mucin type synthetic compounds or it's carrier conjugated compounds. The present invention further concerns the use of non-mucin type synthetic compounds or it's carrier conjugated compounds for preparation of monoclonal antibodies, human immunodeficiency virus (HIV) agents, antitumor agents and immunostimulants.

2.Current Technology

Mucin type antigens such as Tn (GalNAc α 1 \rightarrow O-Ser/Thr), TF (Gal β 1 \rightarrow 3GalNAc α 1 \rightarrow O-Ser/Thr), STn (NeuAc α 2 \rightarrow 6GalNAc 1 \rightarrow O-Ser/Thr) as shown in following below figure, are highly expressed in tumor tissues, and appearance in the normal tissues are restricted. (G.F.Springer, J. Natl, Cancer Inst., 1975, 54, 335., S.Hakomori, Advanced in Cancer Research, 1989, 52, 257)

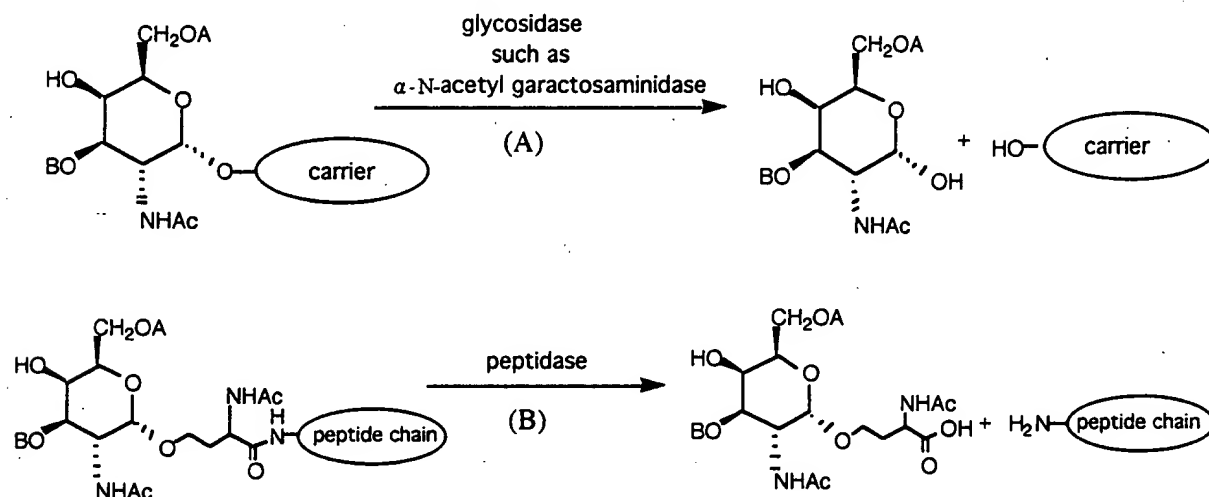


Recently, Tn and STn epitopes, were found on the gp120 as human immunodeficiency virus (HIV) specific glycoprotein. (Hanse, J. E., J. Virol.; 1990, 64, 2833., J. Virol., 1991, 65, 6416.; Arch. Virol., 1992, 126, 11.) And it was reported that the monoclonal antibodies for the O-linked oligosaccharide block HIV infections. (Hanse, J. E., J. Virol.; 1990, 64, 2833.; Kumar A., Virology, 2000, 274, 149.)

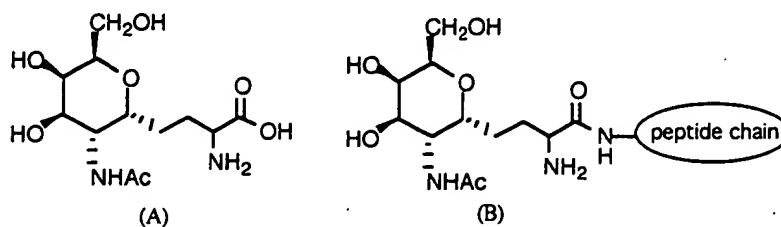
The administration of mucin type tumor antigens and/or attached to pharmaceutically acceptable

carriers are expected as a specific immunotherapy for cancer and HIV. Wherein these carriers are pharmaceutically acceptable proteins such as albumine (ALB), Keyhole limpet hemocyanin (KLH), BCG, or synthetic compounds such as palmitoyl derivatives, aromatic compounds, aliphatic compounds, alkyl, aminoalkyl, peptide and peptoid, which can obtain induction of immune response. (S. J. Danishefsky, J. Am. Chem. Soc. 1998, 120, 12474.; G. Ragupathi, Glycoconjugate J., 1998, 15, 217.; B. M. Sandmeier, J. Immunotherapy, 1999, 22(1), 54.; A. Singhal, Cancer Res., 1991, 51, 1406.; T. Shimizu, 1987, 55, 2287-2289.)

However, above mentioned mucine type antigens-carrier have O-glycoside linkage between sugar and carrier moiety. Therefore, considering about their metabolic stability and immunogenicity, these O-glycoside linkage are susceptible to hydrolyze by glycosidase such as N-acetyl galactosaminidase (EC 3.2.1.49) (Eq A), or hydrolyze of peptide bond by peptidase, as shown in following equation (Eq B), and their activities are anticipated to decrease or attenuate.



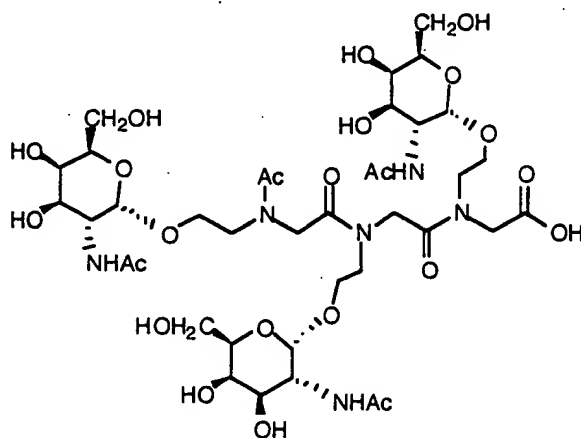
On the other hand, Beau et al have been synthesized the C-glycosides (GalNAc α 1 \rightarrow CH₂-Ser) that have the carbon atom instead of the oxygen atom which connect with serine and N-acetyl galactosamine as a metabolically stable Tn antigen, shown in following equation (A). This Tn antigen is stable against glycosidase such as N-acetyl galactosaminidase (Beau et al, J. C. S. Chem. Commun., 1998, 955.). But, when these C-glycosides are attached to peptides, these compounds could be hydrolyzed by peptidases, and their stabilities are not satisfactory in a living body.



And Roy et al have been synthesized the glycopeptoids as the metabolic stable mimic of carbohydrate

antigen, which was attached to peptoid that is metabolically stable against hydrolysis by peptidase, as shown in following equation (Tetrahedron Lett., 1997, 38, 3487.). But, his reported compounds are thought to be unstable, because it is also expected to be hydrolyzed by glycosidases such as N-acetyl galactosaminidase.

Beau and Roy have not reported to pharmacological activities of Tn antigen linked to carrier proteins.



As mentioned above, the compound, prepared from a coupling of the preceded naturally occurring mucin type antigen and the carrier, is hydrolyzed at it's glycoside bond by glycosidase which exists widely in a living body and at it's peptide bond by peptidase. Therefore it is expected to obtain an insufficient effect.

3.The Subject of Invention

Under these situations, this invention was attained to solve these problems. The purposes of this invention are to prepare the non-mucin type synthetic compounds-carrier conjugated compounds which are stable to hydrolyze against both enzymes, glycosidase and peptidase.

And to prepare non-mucin type synthetic compounds-carrier conjugated compounds which have the ability of specific reactivity to induce immune response for cancer and HIV and have excellent active immunization activities.

And to prepare non-mucin type synthetic compounds-carrier conjugated compounds which are able to obtain selective monoclonal antibodies for cancer and HIV.

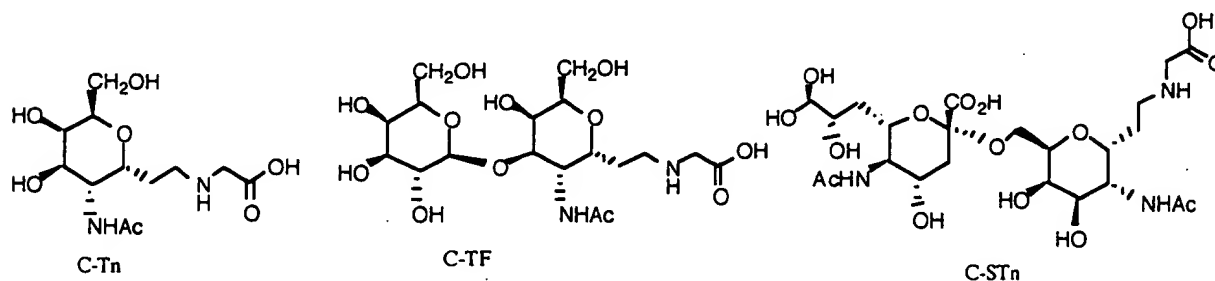
And to prepare anti-tumor agents, anti-HIV agents and immunostimulants which contain this non-mucin type synthetic compounds-carrier conjugated compounds as active ingredients.

And the method to prepare N-acetyl galactosamine in cost effectively, starting material of non-mucin type synthetic compounds-carrier conjugated compounds, is the another object of present invention.

4. A Solution of the Problem

Under these backgrounds, we maked effort to consider to prepare the metabolic stable compounds,

having C-glycoside and peptoid, against to glycosidase and peptidase and the non-mucin compound as shown in following figure, were synthesized for the first time. These compounds are , so to speak, C-glycopeptoid and new compounds.

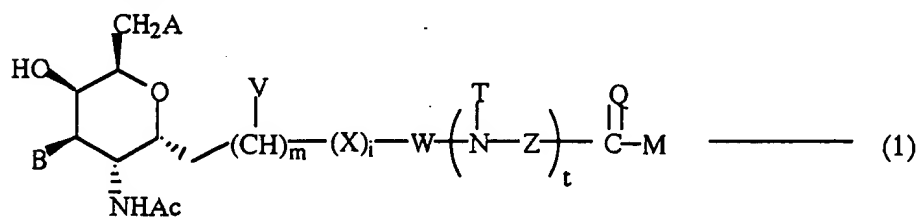


These C-glycopeptoids are more metabolically stable against glycosidase and peptidase than known vaccines. Furthermore, they are able to show their effects for longer time and to store for long term at room temperature. When these novel C-glycopeptoids which are attached to pharmaceutically acceptable carrier proteins, these compounds are more metabolically stable in a living body than known vaccines against glycosidase and peptidase and also expected to show the excellent active immunization activities. It is expected that these novel C-glycopeptoids have potent passive immunogenicities for cancer and HIV. Monoclonal antibodies which were prepared using these compounds are expected to have activities for cancer therapy as a possible immune response. And these compounds have antitumor, anti HIV activity and immunopotentiality.

As a result of investigation of novel compounds as anti HIV agents and immunostimulant, the antibodies which was prepared using novel compound have excellent antitumor, anti HIV activity and immunostimulation as shown in general formula (1).

DETAILED DESCRIPTION OF THE INVENTION

A compound of the general formula (1),



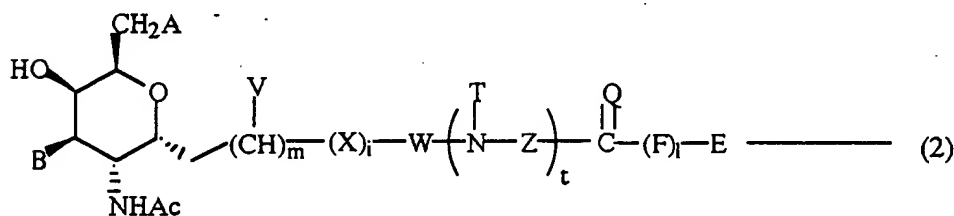
wherein A represents OH or sialic acid and/or its derivatives, and B represents OH or galactose and/or its derivatives; T represents H or protecting groups of amine; M represents H or OH; X represents oxygen atom, -NH- or S(O)_z (where z is 0, 1 or 2); Q is H or oxygen atom; V represents lower alkyl or H; W is straight or branched alkylene groups from 0 to 5; Z is straight or branched alkylene groups from 1 to 5; i, m, and t is 0 or 1;

non-mucin type synthetic compounds or its carrier conjugated compounds, which have above mentioned compounds as a core structure of antigen.

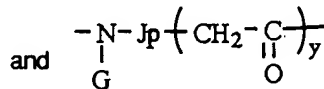
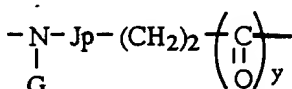
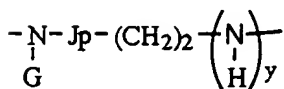
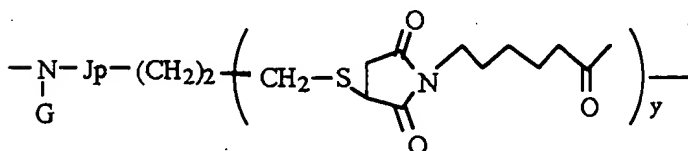
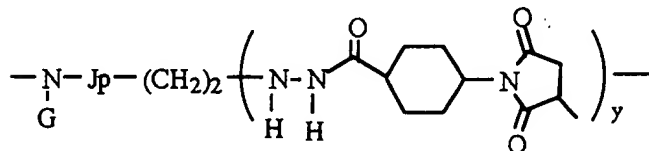
In the explanation of T, protecting groups of amin are alkyl, acetyl, t-butyloxycarbonyl, benzyloxycarbonyl group, and others. W is straight or branched alkylene groups from 0 to 5. And general formula (1) is called non-mucin compound in this invention.

Compound (1) have immunopotentiality and non-mucin type synthetic compounds or its carrier conjugated compounds can be prepared from compound (1) or 2-5 clustered of compound (1) linked with the synthetic compounds such as palmitoyl derivatives which can obtain inductions of immune response.

A compound of the general formula (2),

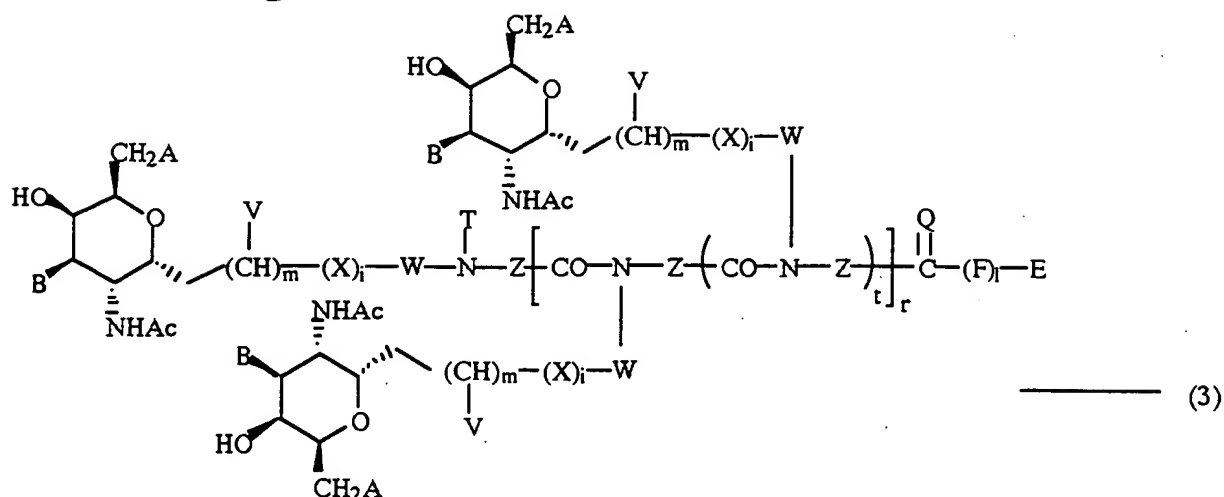


wherein A, B, T, X, Q, V, W, Z, i, m, and t have above-mentioned meanings; E represents pharmaceutically acceptable carrier compounds; l is 0 or 1; F is showed followings,



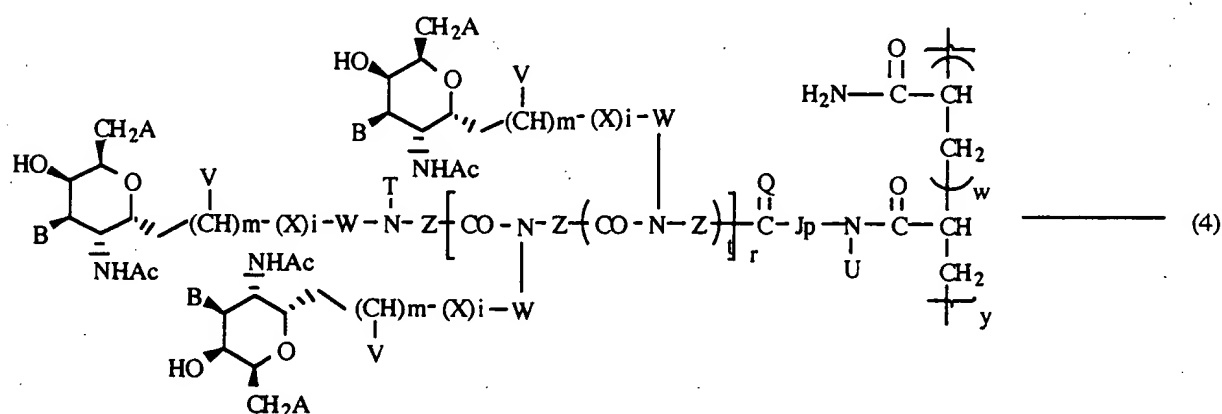
wherein J is $\text{---CH}_2\text{CH}_2\text{X---}$ or $\text{---N(L)---CH}_2\text{CO---}$ (where X have above-mentioned meanings; L is H or lower alkyl); G is H or lower alkyl; p is 0 to 3; y is 0 or 1;

non-mucin type synthetic compounds or its carrier conjugated compounds, which have above compounds as a core structure of antigen.



wherein A, B, T, X, Q, V, W, Z, i, m, t, E, , and l have above-mentioned meanings; r is from 1 to 4; non-mucin type synthetic compounds or it's carrier conjugated compounds, which have above compounds as a core structure of antigen.

. A compound of the general formula (4),



wherein A, B, T, X, Q, V, W, Z, J, i, m, t, p, and r, have above-mentioned meanings; U represents H or lower alkyl; w is 0 to 50; y is 1 or 50.

In the explanation of E, pharmaceutically acceptable proteins are such as albumine (ALB), Keyhole limpet hemocyanin (KLH), BCG, or synthetic compounds such as palmitoyl derivatives, aromatic compounds, aliphatic compounds, alkyl, aminoalkyl, peptide and peptoid, which can obtain induction of immune response.

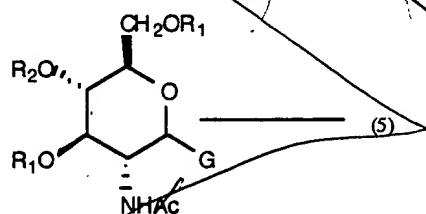
Non-mucin type synthetic compounds or it's carrier conjugated compounds contain general formula (1) as a core structure. These compounds are able to apply to mammal such as human, and these are used as anti-tumor agents and/or anti-HIV agents having immunostimulate activities. These compounds are also used for the preparation of monoclonal antibodies. These novel compounds are

able to elongate of effective time, to decrease of dosage, and to reduce of side effects, further the compounds of the invention are expected to have the potent immunogenicities for cancer and HIV than known vaccines. It is expected that monoclonal antibodies, prepared from this invention, have excellent antitumor and anti HIV activity. Furthermore, when neuraminidase inhibitors such as Zanamivir or Oseltamivir are co-administrated with sialic acid contained compounds in this invention, these sialic acid contained compounds are expected to be more stable in a living body.

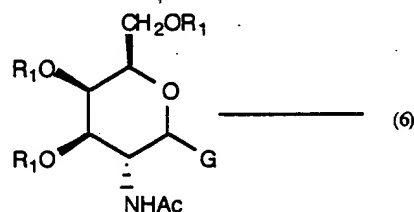
N-acetyl galactopyranose moiety in mucin type (O-Tn, O-STn, O-TF) or non-mucin type (C-Tn, C-STn, C-TF) antigens was synthesized from N-acetyl galactosamine that is very expensive as a starting material. On the other hand, N-acetyl glucosamine, isomer of N-acetyl galactosamine at C-4 hydroxy group, is cheaper and readily available. So it is hoped to use cheaper N-acetyl glucosamine as starting material.

Following this invention, N-acetyl galactosamine derivatives can be synthesized from N-acetyl glucosamine via inversion of C-4 hydroxy group.

That is to say the process for the preparation of N-acetyl galactosamine derivatives, general formula (6) can be prepared from the invention of OR₂ group to OR₁ group at C-4 position in N-acetyl glucosamine derivatives, general formula (5).



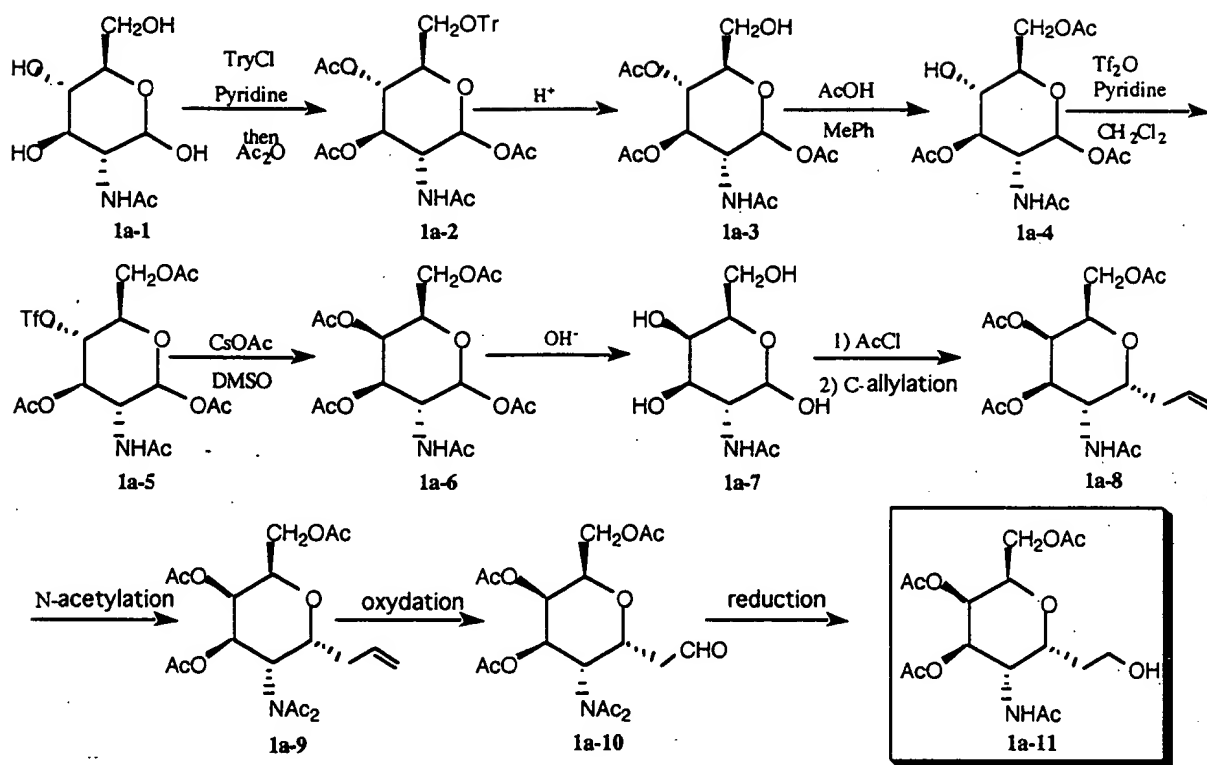
wherein OR₁ is H or a protecting group of a hydroxy group such as acetyl group; R₂ is a leaving group such as tosylate, trifluoromesylate or methansulfonate; G is allyl or protected hydroxyl groups.



Herein we described the method for preparing the key intermediate, galactose derivatives, (1a-11), and also general formula (1).

1) Synthesis of intermediate 1a-11

(i) Route 1-a



The intermediate 1a-11 is synthesized starting from readily available N-acetylglucosamine as shown in route 1-a via inversion of C-4 hydroxy group.

N-acetylglucosamine is selectively protected by trityl ether at C-6 position (B. Helferich et al, Ann., 1920, 450, 219.), followed by acetylation at C-3, 4 and treated with formic acid afford compound 1a-3 (M. Bessodes, Tetrahedron Lett., 1986, 27, 579.).

The 4-hydroxyl intermediate 1a-4 is obtained via acetyl migration of the compound 1a-3 by heating with acetic acid in toluene (D. Chaplin et al, J. Chem. Soc. Perkin Trans. 1, 1992, 235.).

The preparation of 4-hydroxyl derivative is selectively protected as benzoyl or pivaloyl ester at position C-3 and 6 by only one step procedure.

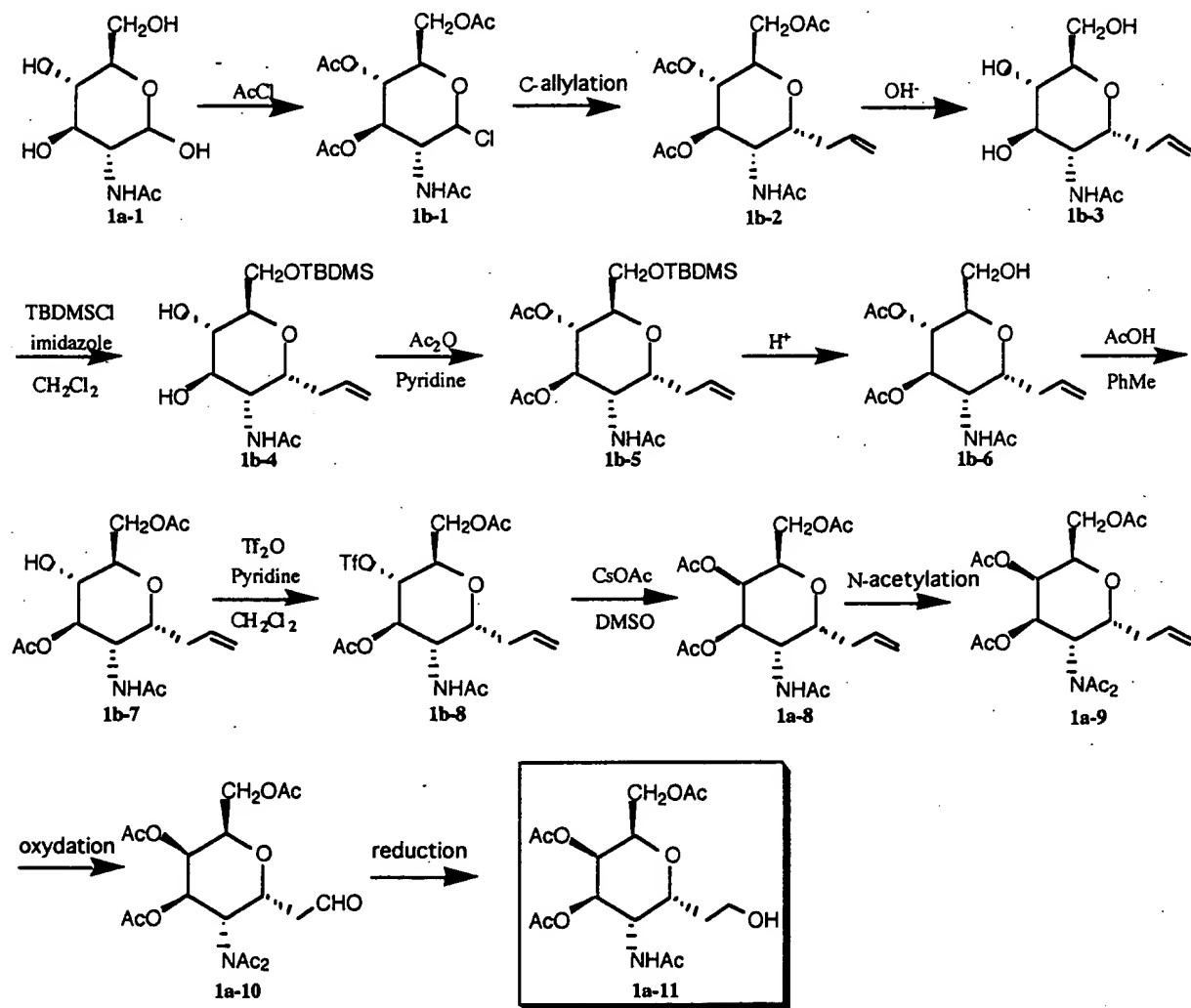
4-hydroxyl group is transformed to triflate 1a-5 and the inversion step is carried out using cesium acetate to give N-acetyl-1,3,4,6-tetra-O-acetyl-D-galactosamine 1a-6.

Methanesulfonyl chloride or p-toluenesulfonyl chloride can be used instead of trifluoromethanesulfonyl chloride. Then compound 1a-6 is deacetylated to N-acetyl-D-galactosamine. And this epimerization at C-4 position was carried out by the procedure of Cipolla et al (Tetrahedron Asymmetry, 2000, 295-303). Allyl group is introduced into compound 1a-7 by Horton's procedure (Carbohydr. Res., 1996, 309, 319-330).

Compound 1a-7 is reacted with acetyl chloride, followed by allylation using allyltributyltin and 2, 2'-azobis isobutyronitril (AIBN) to obtain compound 1a-8. But, the method of allylation is not restricted by this allylation method.

2-Acetamide group is protected as N, N-diacetyl using isopropenyl acetate in the presence of catalytic amount of acid to afford compound 1a-9 (J. Owi. Horton et al, Carbohydr. Res, 1996, 309, 319-330.). Compound 1a-9 is reacted with osmium oxide and NaIO₄ to obtain aldehyde compound 1a-10. Compound 1a-10 is subjected to reduction using sodium brohydride to give compound 1a-11.

(ii) Route 1-b

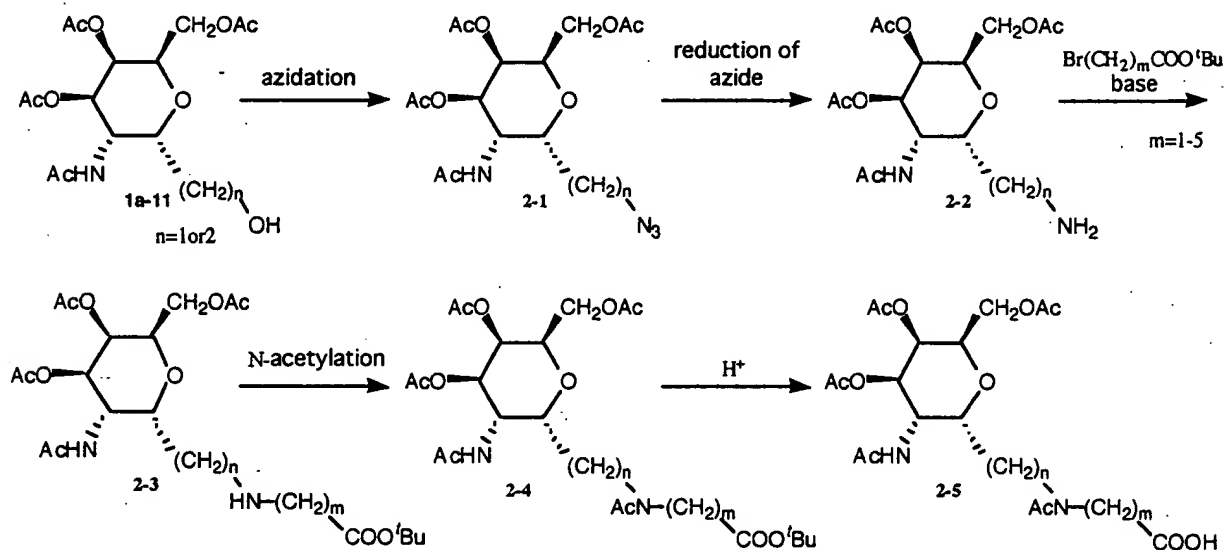


The intermediate 1a-11 is also synthesized starting from N-acetyl-D-glucosamineas shown in route 1-b. The compound 1a-1 is inverted at C-4 hydroxyl group after induction of allyl group (B.A.Roe et al, J. Org. Chem, 1996, 61, 6442-6445.). N-acetyl-D-glucosamine is treated with acetyl chloride followed by C-allylation with allyltributyltin afford compound 1b-2. Compound 1b-2 is deacetylated

with NaOMe to afford compound 1b-3. Then compound 1b-3 is selectively protected as t-butyldimethylsilyl (TBS) ether at C-6 position, followed by acetylation with acetic anhydride under basic conditions to give compound 1b-5. Compound 1b-5 is desilylated by acids and rearrangement to compound 1b-7 by heating with a catalytic amount of acetic acid in toluene.

Synthesis of compound 1a-11 from 4-hydroxyl compound 1b-7 is obtained by similar method as described in route 1-a.

2) Synthesis of compound 2-5: Route 2



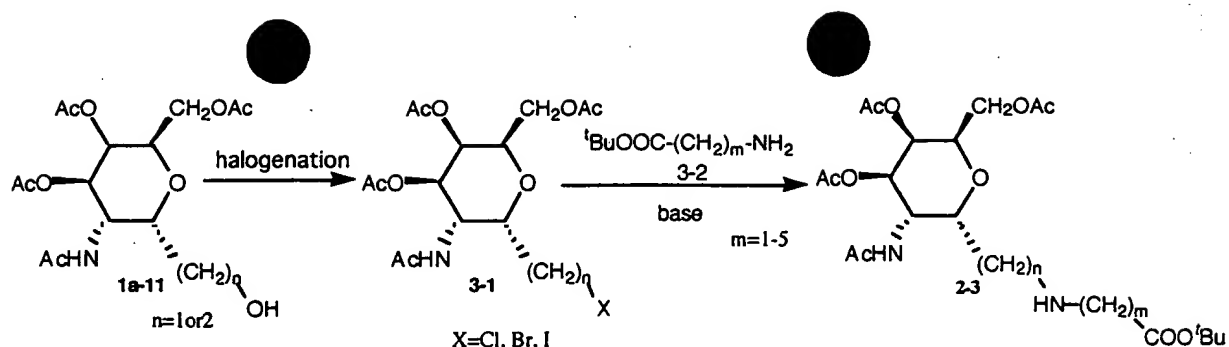
Compound 1a-11 is converted to 2-1 by Mitsunobu reaction (O.Mitsunobu, Synthesis, 1, 1981.).

Then azide group of compound 2-1 is reduced to primary amine ; for example hydrogenation using Pd-C.

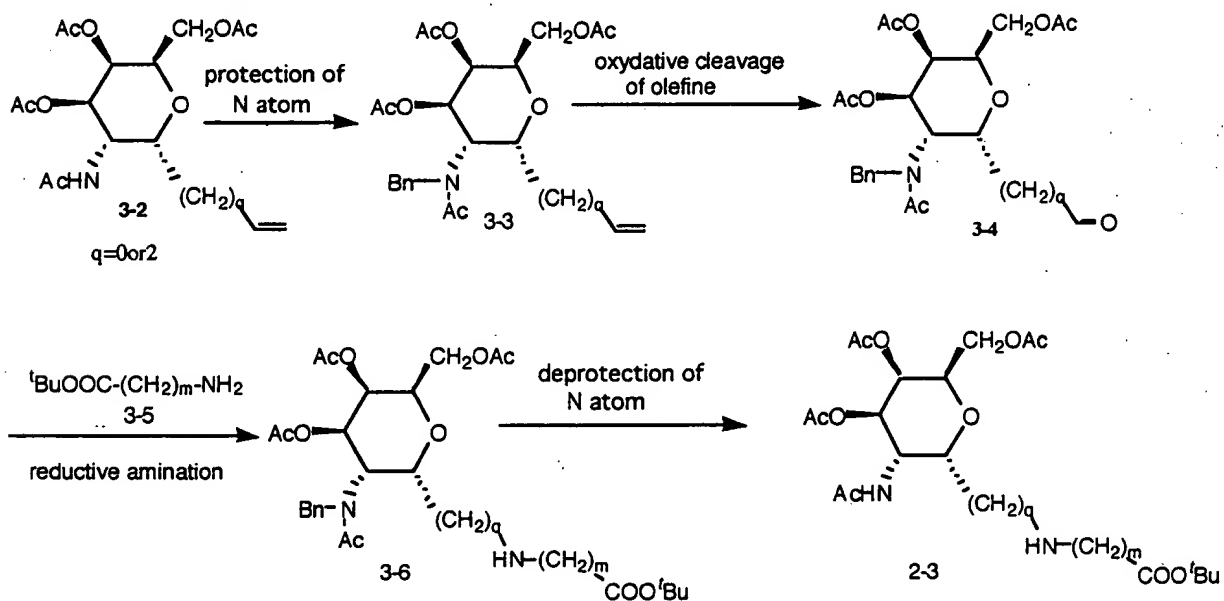
Alkylation of compound 2-2 with haloester, for example butyl bromo acetate, give compound 2-3.

Compound 2-3 is protected as acetamide using acetic anhydride or acetyl chloride. Compound 2-5 was obtained by deprotection of compound 2-4.

3) Synthesis of compound 2-3: Route 3

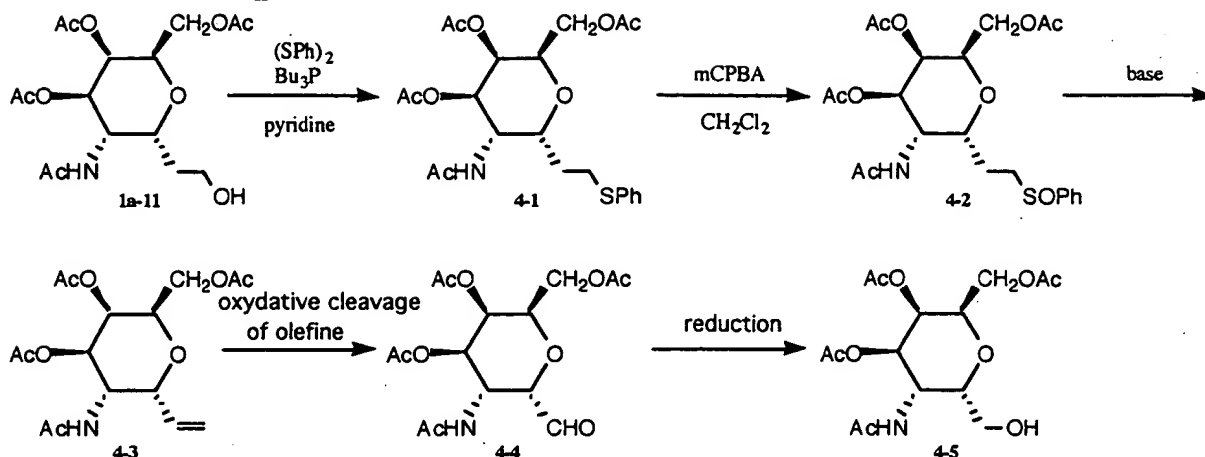


The hydroxyl group in compound 1a-11 is converted to leaving group such as halogen, followed by coupling with compound 3-2 in the presence of base give compound 2-3. But, the leaving group is not restricted to halogens.



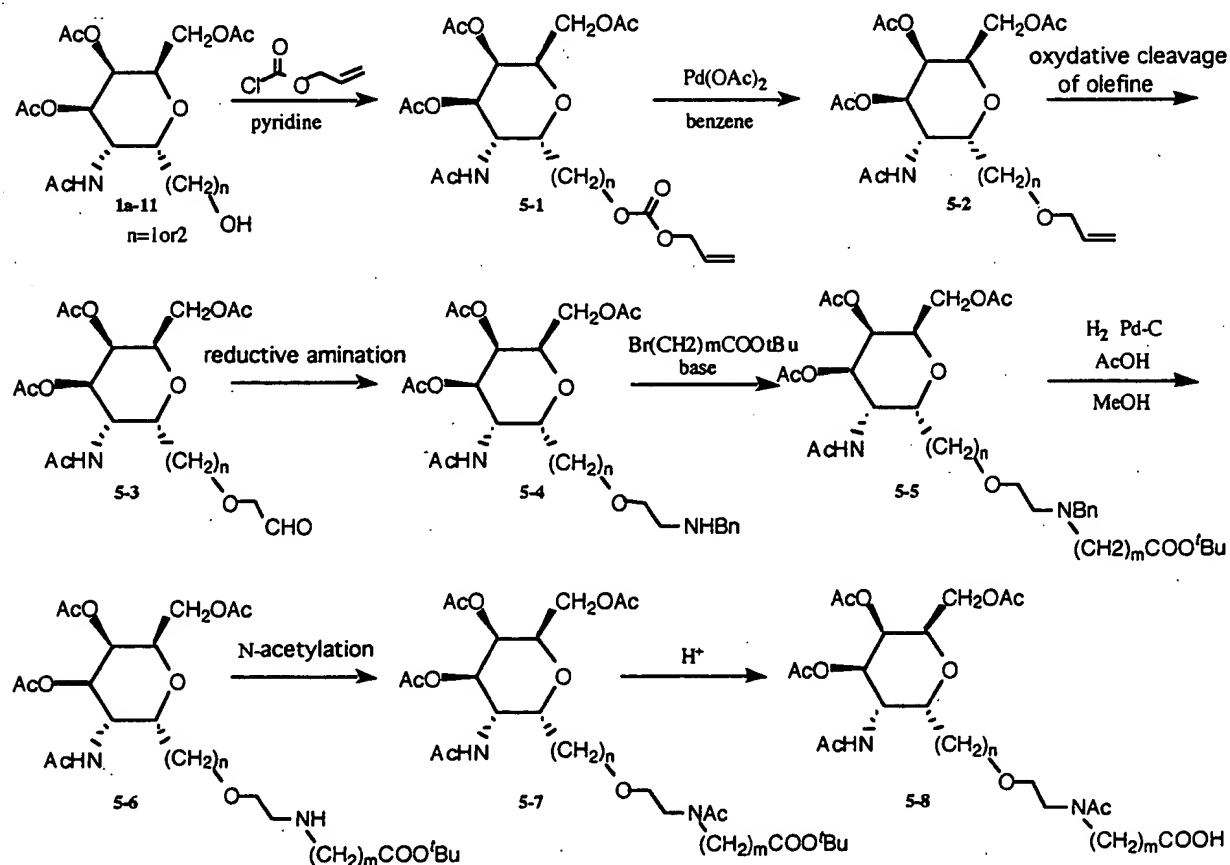
Acetamide group in compound 3-2 is protected as suitable protecting group, for example benzylamide, followed by oxydative-cleavage of olefine to provide aldehyde 3-4. Reductive amination with compound 3-5, followed by deprotection give compound 2-3.

4) Synthesis of compound 4-5: Route 4



Compound 1a-11 is reacted with diphenyldisulfide to give compound 4-1, followed by oxidation using *m*-chloroperbenzoic acid to give compound 4-2. Then, heating in the presence of amine give olefin compound 4-3. The compound 4-3 is oxidized, followed by reduction give compound 5-8 as described route 1.

5) Synthesis of compound 5-8: Route 5.

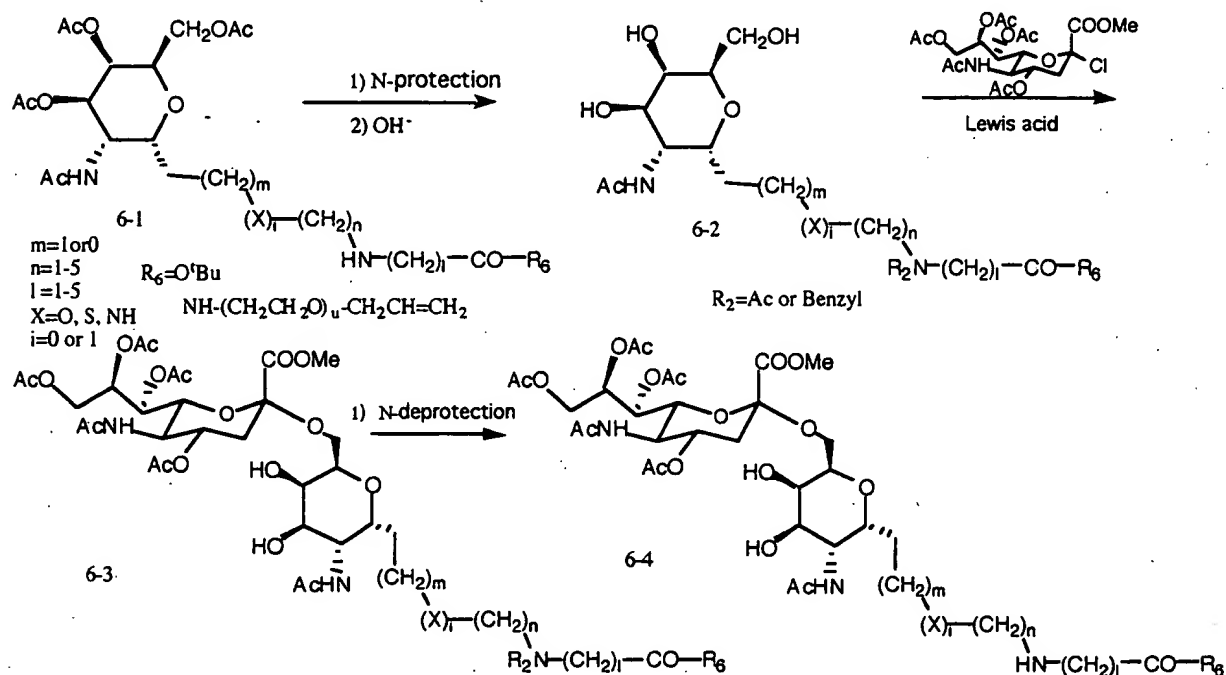


Allyl group is introduced into compound 1a-11 by the method of Curibe et al (Tetrahedron Lett.,

1981, 22, 3591-94). Ozonolysis or oxidative cleavage of compound 5-2 afford compound 5-3 by OsO_4 . The reductive amination of compound 5-3 using amine, for example benzylamine, give compound 5-4. The compound 5-4 is coupled with haloester (for example butyl bromoacetate) provide compound 5-5. Then amino group of the compound 5-5 is deprotected by hydrogenation, followed by acetylation and debutylation give compound 5-8.

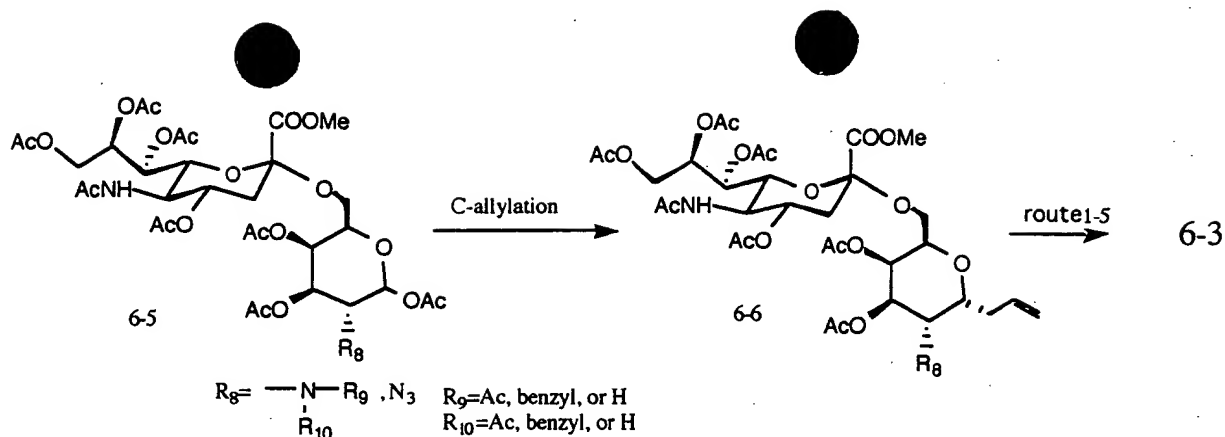
6) Synthesis of sialyl acid derivatives: Route 6

(i) Route 6-a



The protection of amino group, followed by deacetylation give compound 6-2. The compound 6-2 is glycosylated with sialic acid derivatives by Danishefsky's procedure (J. Am. Chem. Soc., 1999, 121, 2662-2673.). And, leaving group in this reaction is not restricted to halogens. Obtained compound 6-3 can be converted to compound 6-4 as a intermediate of cluster.

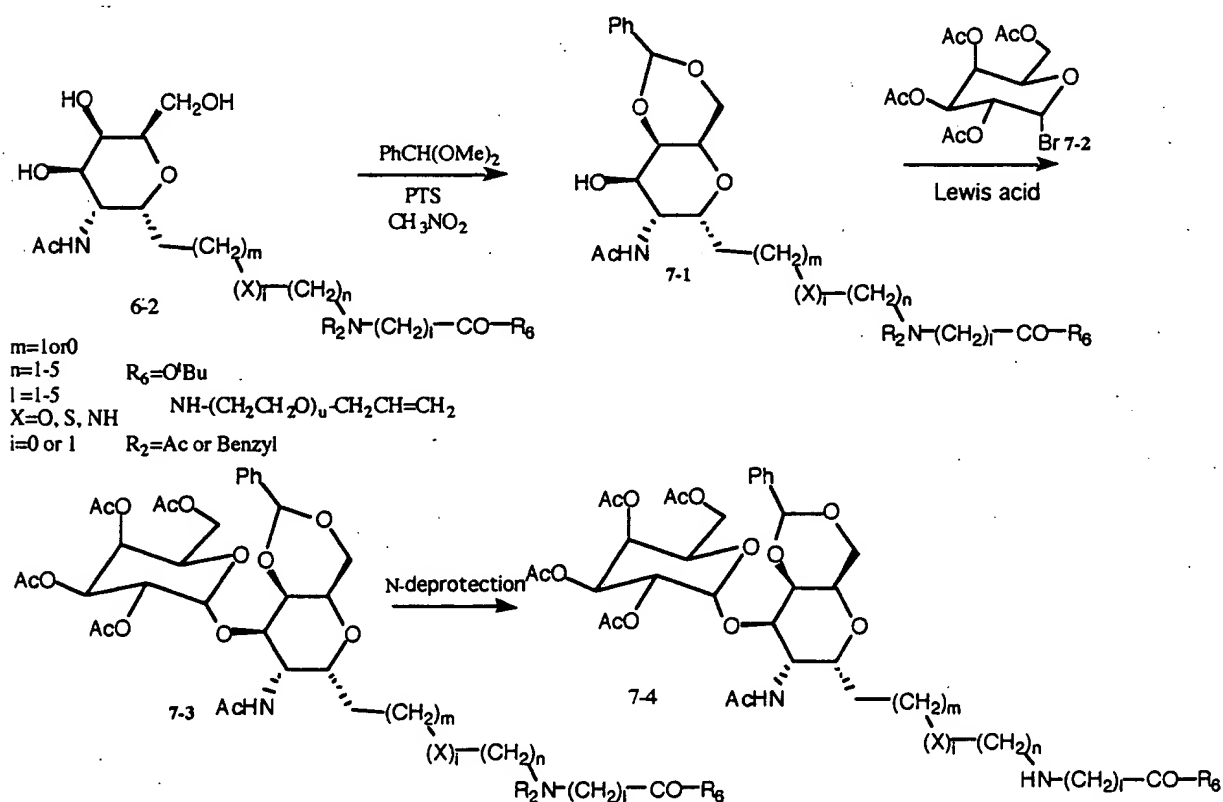
(ii) Route 6-b



The α -C-glycoside 6-6 is obtained by C-allylation of compound 6-5. Synthesis of compound 6-3 from compound 6-6 is similar to described route 1-5. And this reaction also can be proceeded using sialyl transferase and sialic acid derivatives (C.Pauison et al. J. Am. Chem. Soc., 1990, 112, 9308-9309).

7) Synthesis of galactose derivatives: Route 7

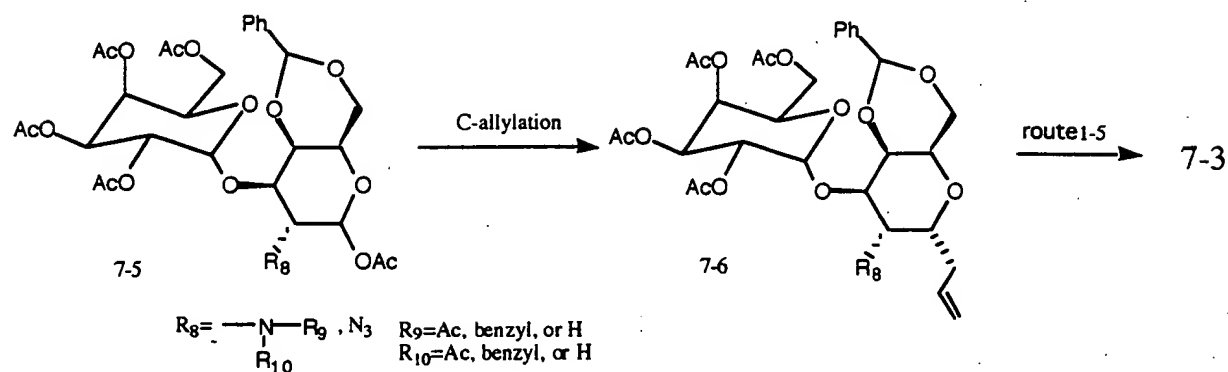
(i) Route 7-a



Compound 7-1 is obtained by hydroxyl group protection of compound 6-2. The compound 7-1 is glycosilated with acetobromo galactose to give compound 7-3. And, the leaving groups in this

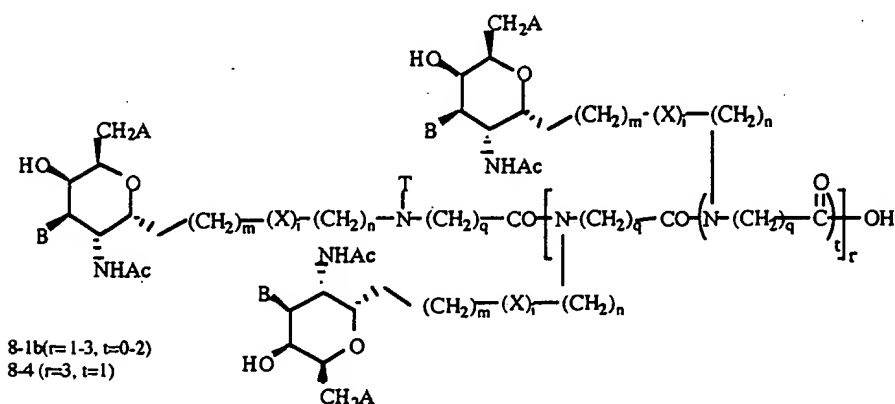
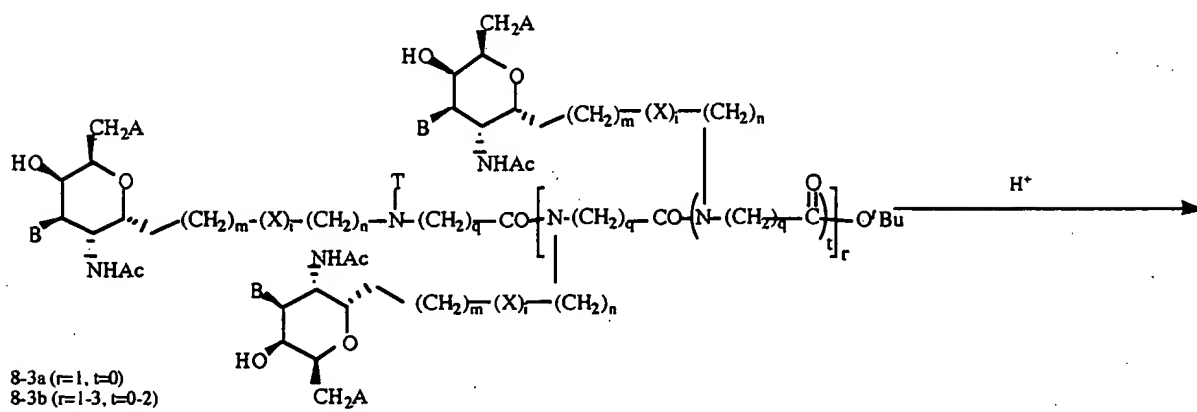
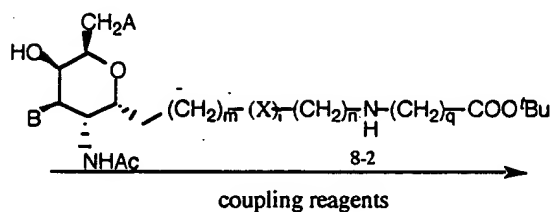
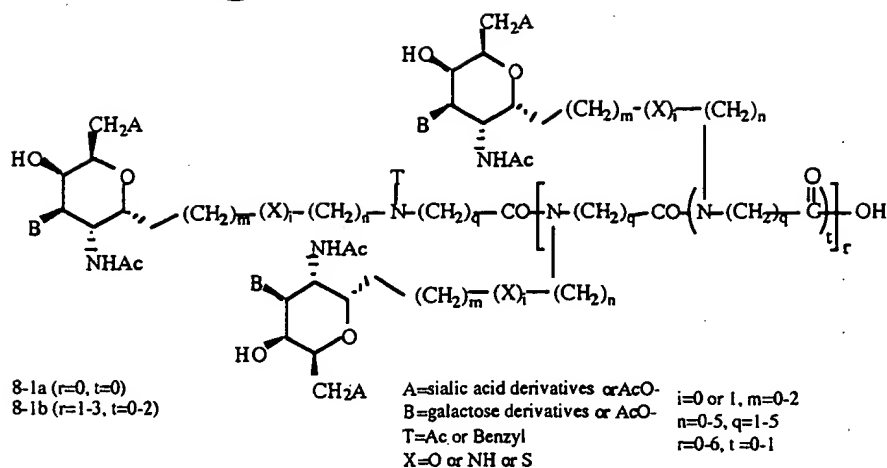
reaction is not restricted to halogen. Compound 7-3 can be converted to compound 7-4 as an intermediate of cluster.

(ii) Route 7-b



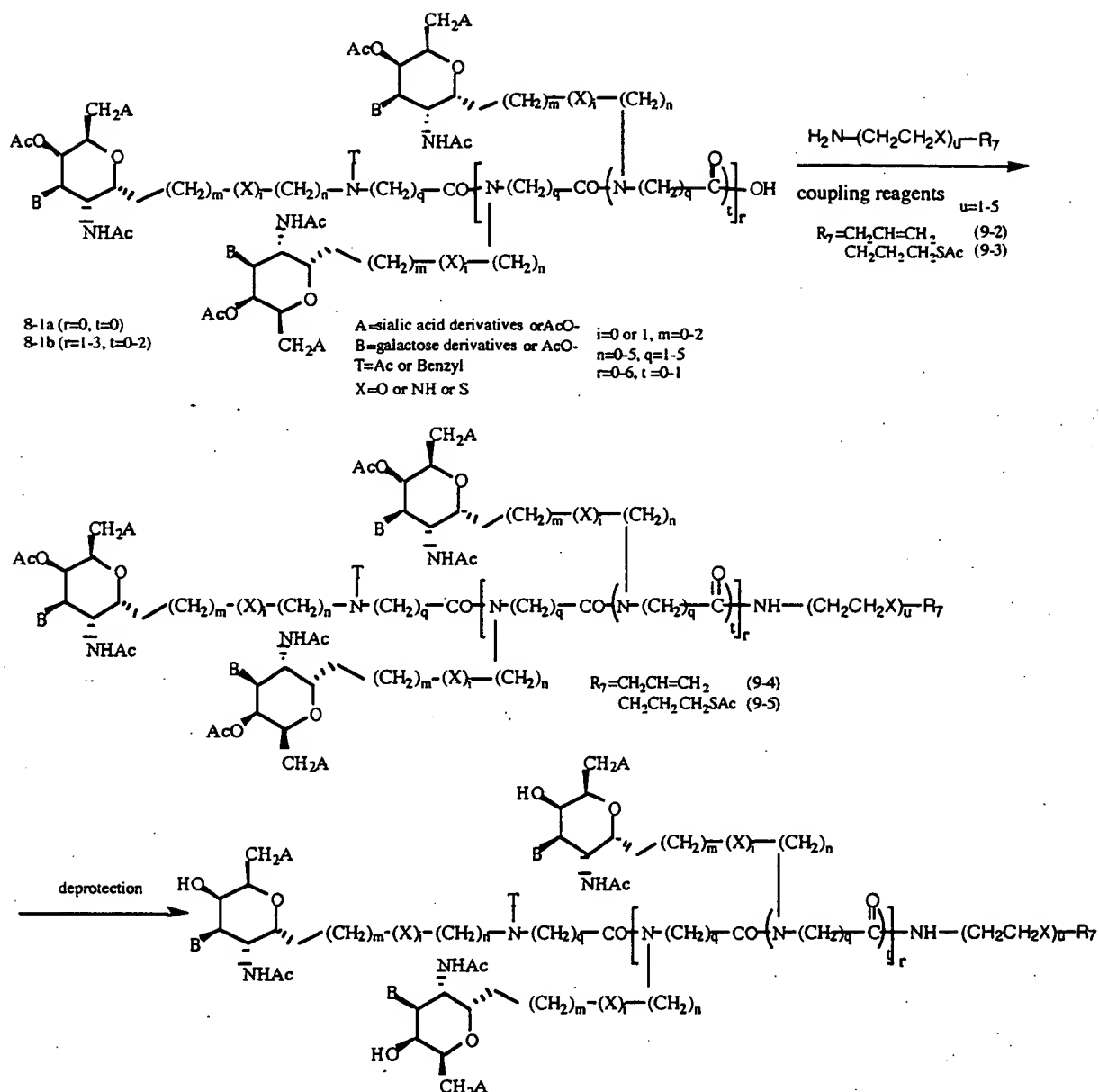
The α -C-glycoside 7-6 is obtained by C-allylation of compound 7-5. Synthesis of compound 7-3 from compound 7-6 is similar to described route 1-5. And this reaction also can be proceeded using sialyl transferase and sialic acid derivatives (C. Pauison et al. J. Am. Chem. Soc., 1990, 112, 9308-9309).

8) Synthesis of culster derivatives: Route 8



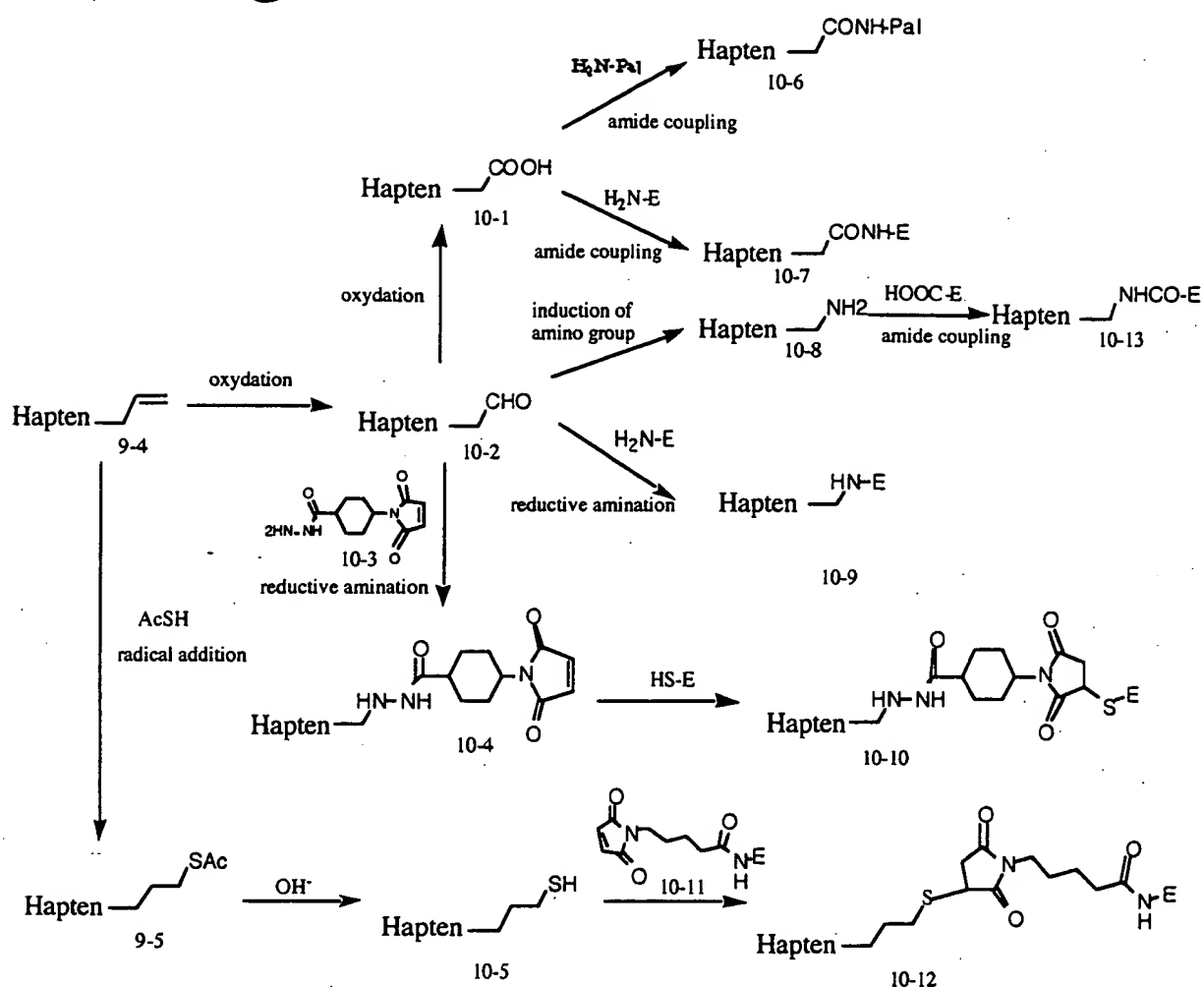
Compound 8-3 is obtained by coupling compound 8-1a with compound 8-2 by P.Roy's procedure (Tetrahedron Lett., 1997, 38, 13478-13490.). Compound 8-1b is synthesized by deprotection of compound 8-3, followed by coupling with compound 8-2 afford compound 8-3b. Compound 8-1b, 4 is obtained by deprotection of ester in compound 8-3a, b.

9) Coupling linker with haptens: Route 9



Compound 9-4 and 9-5 are prepared from coupling linker 9-2 or 9-3 with compound 9-1 respectively (P. Roy et al, Tetrahedron Lett., 1997, 38, 3478-3490). And this coupling can be proceeded used to other reagents such as N, N-dicyclocarbodiimide, Danishefsky's procedure (J. Am. Chem. Soc., 1998, 120, 12474-12485) or 2-isobutyl-1-isobutoxycarbonyl-1,2-dihydroquinoline. Compound 9-7 is obtained by deprotection of hydroxy group.

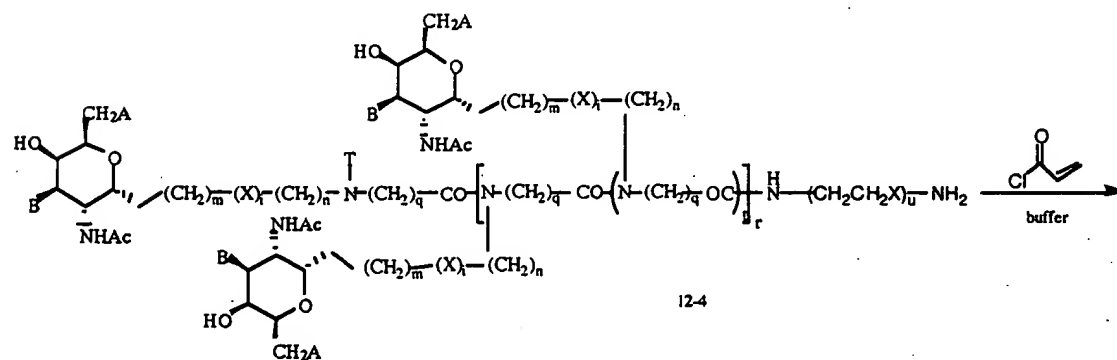
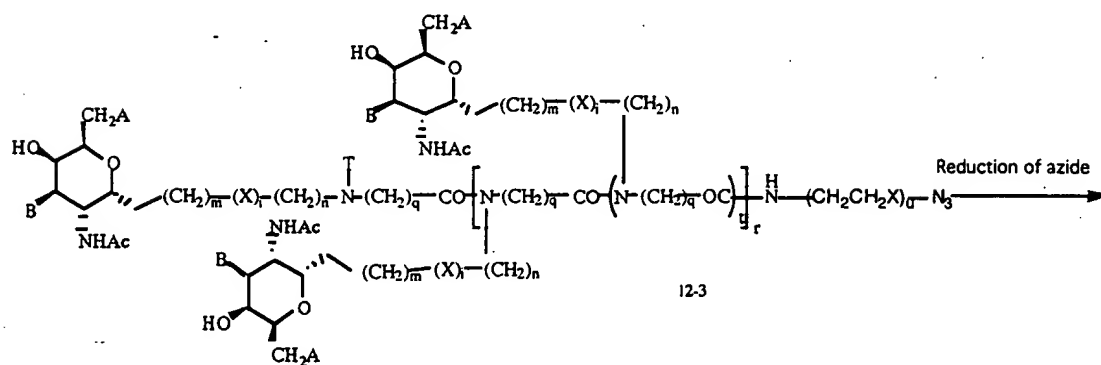
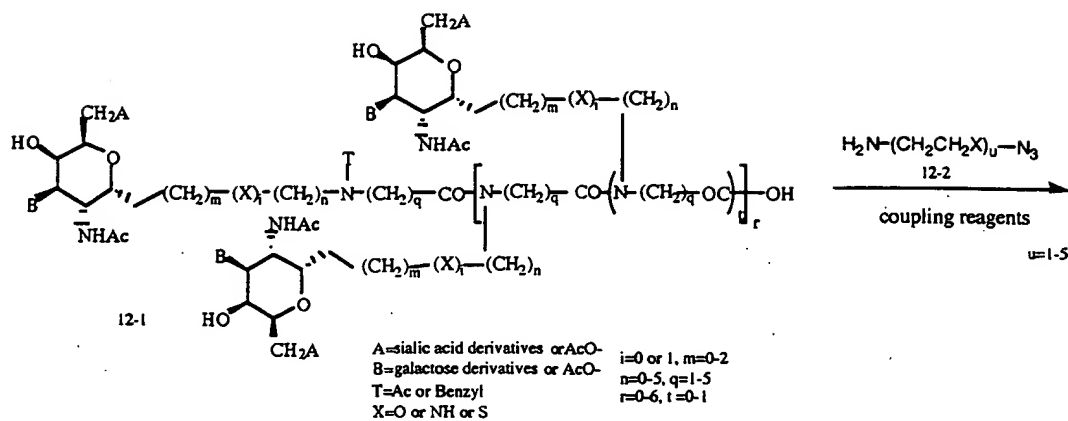
10) Coupling carrier with hapten : Route 10

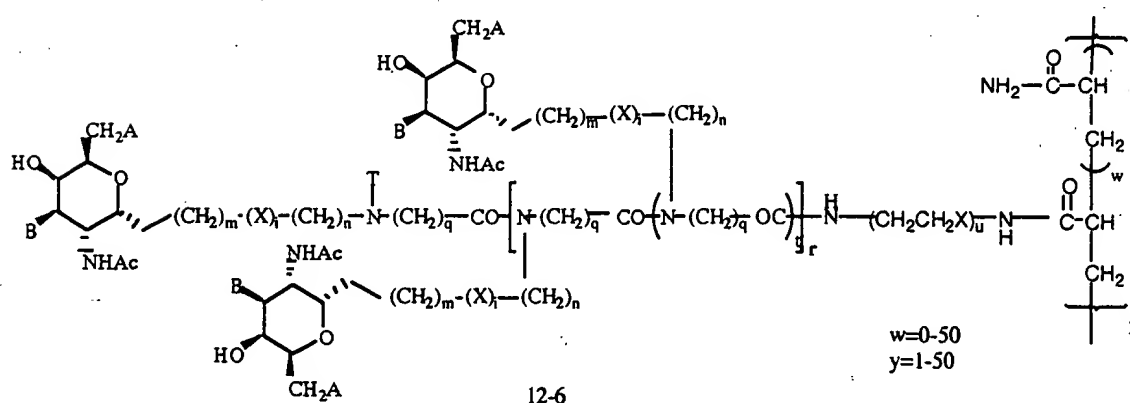
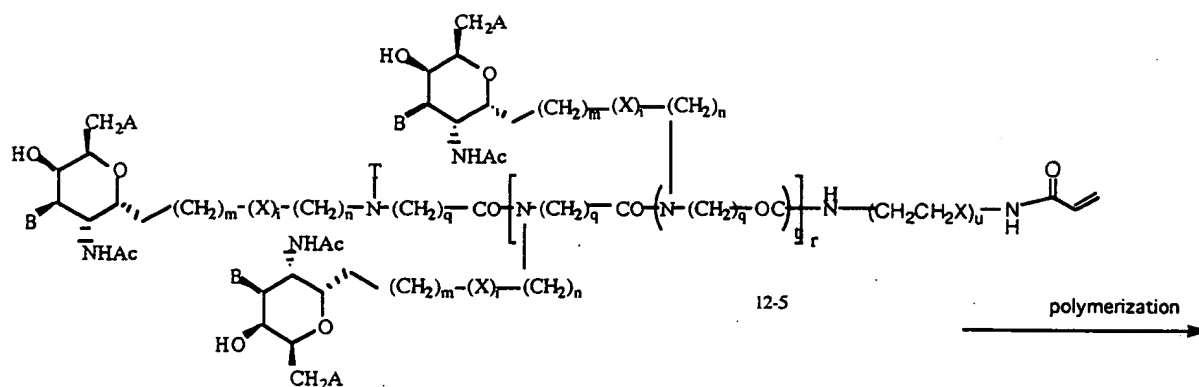


Compound 9-4 is oxidized to aldehyde 10-2, followed by reductive amination with protein using sodium cyanoborohydride in phosphate buffer (pH7.2) afford compound 10-9 by Livingston's method (Glycoconjugate Journal, 1998, 115, 217-221.). Compound 9-4 is also coupled with protein by Slovin's method (Proc. Nat. Acad. Sci. USA, 1999, 96, 5710.). Compound 9-4 is converted to maleimide compound 10-4, followed by coupling with thiol group in protein or compound 9-4 is converted to thioacetate compound 9-5. Compound 9-5 is coupled with maleimidated protein by Khono's method (J. Clin. Lab. Anal., 1999, 10, 91.) afford compound 10-12.

Compound 10-7 is obtained by coupling carboxylic group in compound 10-1 with amino groups. Compound 10-13 is obtained by coupling aminogroup in compound 10-8 with carboxylic groups in protein. And Coupling compound 10-1 with palmitoyl derivatives by Danishefsky's method (J. Am. Chem. Soc., 1999, 121, 2662-2673.) give compound 10-6.

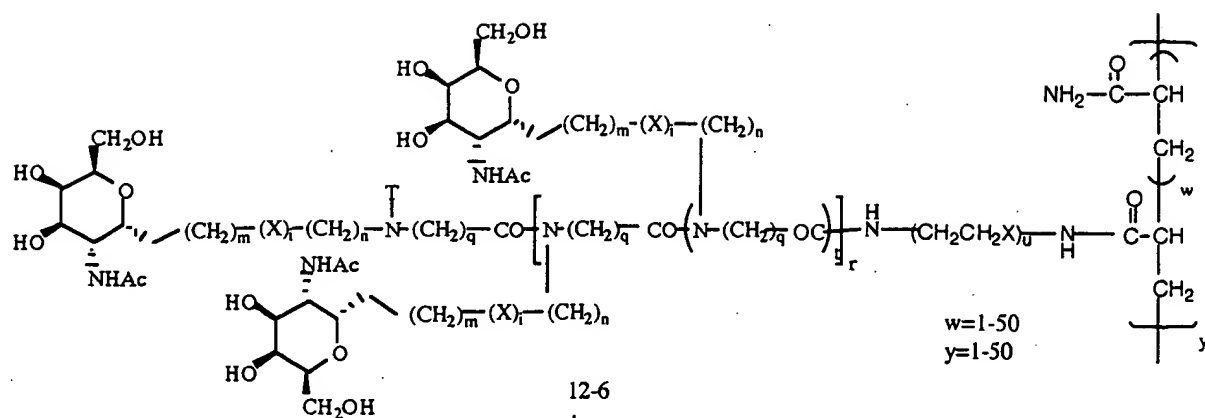
11) Synthesis of polymer derivatives: Route 11-a





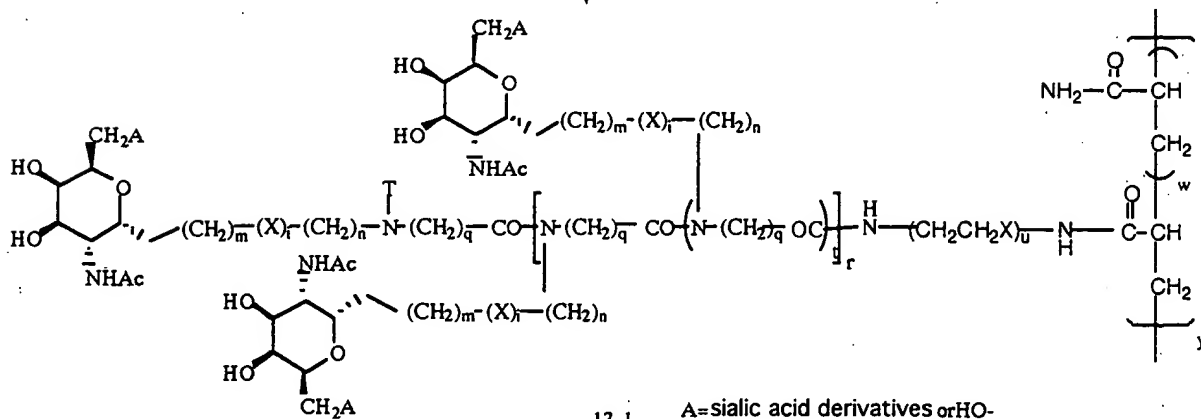
To couple a linker 12-2 with carboxylic acid compound 12-1 afford compound 12-3. Compound 12-3 is reduced to compound 12-4, followed by condensation with acryloyl chloride and afforded compound 12-5. Compound 12-5 is converted to compound 12-6 by the method of K.Eklind et al (J.Carbohydrate Chem., 1996, 15, 1161) or J.Domb et al (J.Med.Chem., 2000, 43, 2591). The method of polymerization is not restricted to this method.

(ii) Route 11-b



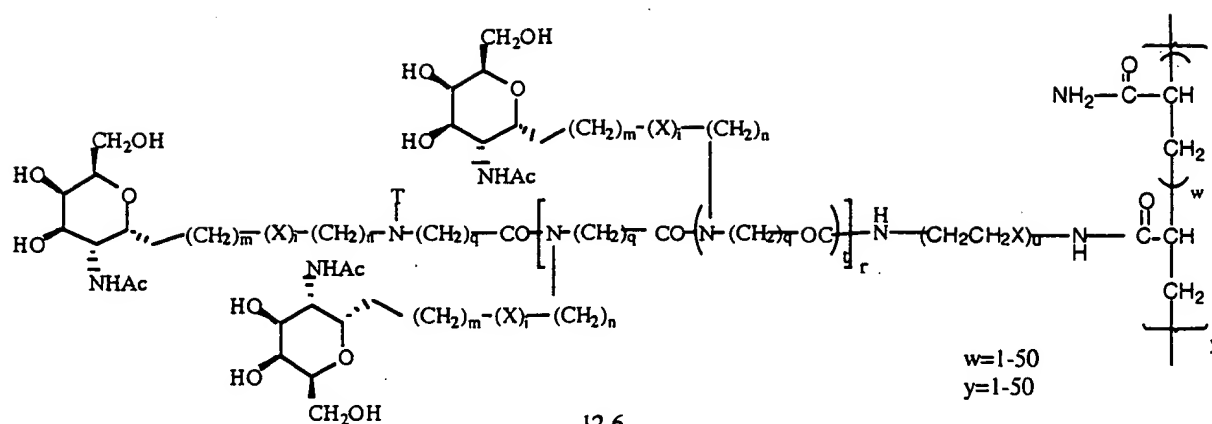
12-6

CMP-Neu-5Ac
2,6-sialyltransferase



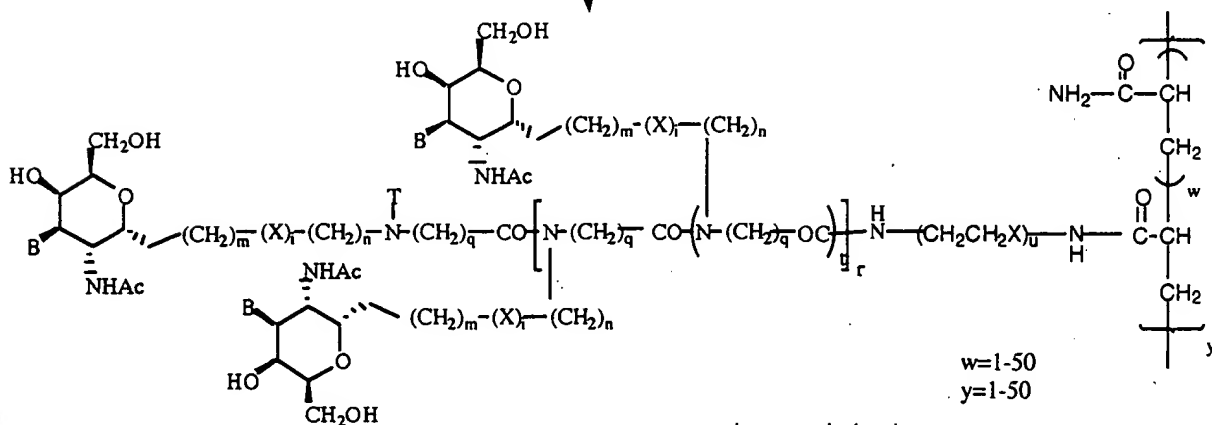
Synthesis of sialic acid derivative 13-1, having a polymer, can be achieved using CMP-Neu-5Ac and sialyltransferase from compound 12-6 (C. Pauison, J. Am. Chem. Soc., 1990, 112, 9308-9309).

(iii) Route 11-c



12-6

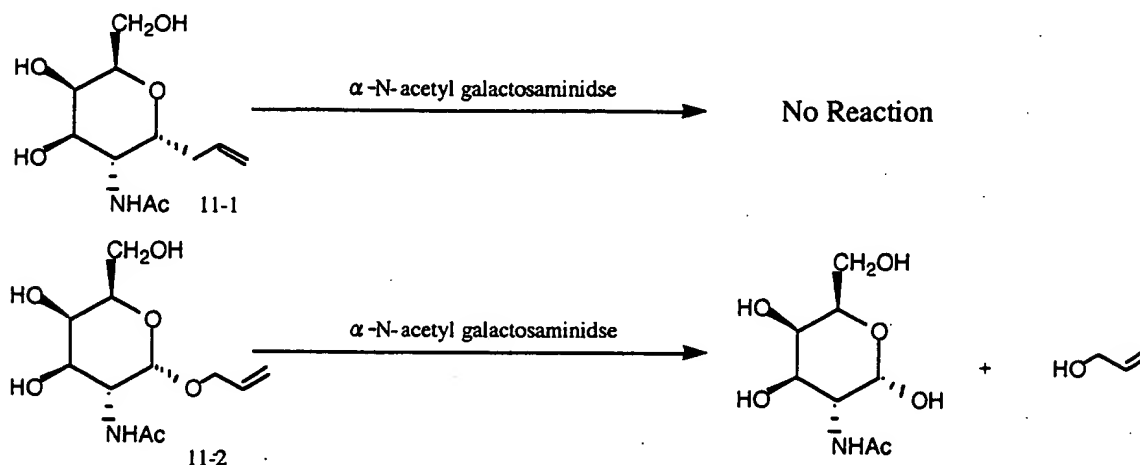
PNP-Gal
Galactosidase



14-1 B=galactose derivatives or HO-

Synthesis of galactose derivative 14-1 can be achieved from compound 12-6 using galactosidase and galactose derivatives PNG-Gal after polymerization (C. Pauison, J. Am. Chem. Soc., 1990, 112, 9308-9309).

The Biological stability for glycosidase such as N-acetyl galactosaminidase was tested using allyl derivatives (11-1, 11-2) by Mark von Itzstein's method (Org. Leyy., 1999, 443-446).



enzyme ; α -N-acetyl galactosaminidase

0.32unit (1.69 unit/ml 0.1%BSA containing 0.5M sodium citrate buffer)

solvent ; citric acid buffer (pD=3) 0.6ml

temperature ; 35°C

procedure ; Substrate (2mg) was dissolved in citric acid buffer (0.6ml) and α -N-acetyl galactosaminidase (0.32 unit) was added. NMR spectrum was determined in every constant time.

Results of this test , Substrate persistence, were shown in table 29.

Table 29

substrate \ time	2	4	6	8	10	12	18	24
11-2	89	79	68	57	50	45	40	22
11-1	100	100	100	100	100	100	100	100

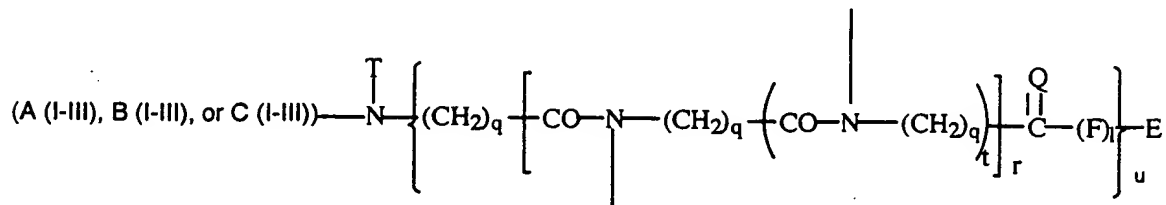
From above results, evidently, 78% of O-glycoside linkage allyl ether (11-2) that have was hydrolyzed after 24 h. As it is expected, compound 11-1, replaced ether bond to C-C bond, was unaffected by enzyme, degradation was not observed after 24h.

This result show the C-glycoside is metabolic and catabolic more stable than O-glycoside.

One or more than two medicinal compounds can be contained compounds that of described general formula (I) in the present invention as active ingredients. And the general formula (1) can be administered to human. And monoclonal antibodies of general formula (I) can be administered to human. The compounds with the invention is concerned may be prepared for administration by any route consistent with their pharmacokinetic propaties. And the compound itself and/or it can be formed injections, powders, granules, tablets, capsules, troches, dry-syrups, lipozome preparations and others. The appropriate dose and dosage times, that of the compound of the present invention, must be determined by the conditions of patient, age, body weight etc.

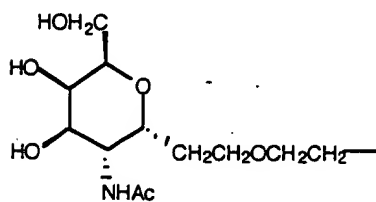
The compounds are exemplified as follow, but the invention is not limited to these compounds.

(A (I-III), B (I-III), or C (I-III))

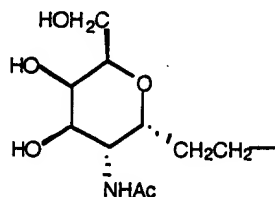


(A (I-III), B (I-III), or C (I-III))

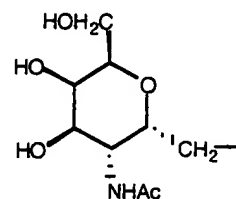
(wherein T, E, F, Q, t, l, and r have above-mentioned meaning; q represents 0 to 5; u is 0 or 1.)
A (I), A (II), A (III), B (I), B (II), B (III), C (I), C (II), and C (III) in above equation and table 1-28 were shown in bellow aquation



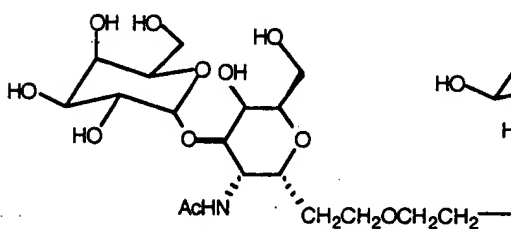
A(I)



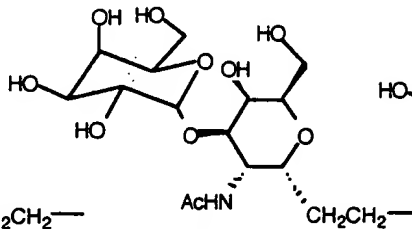
A(II)



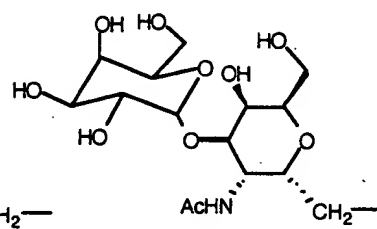
A(III)



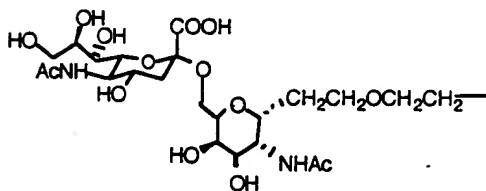
B(I)



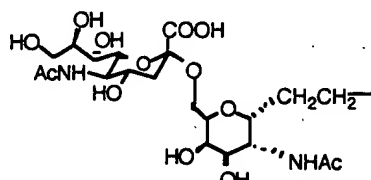
B(II)



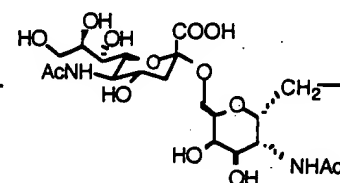
B(III)



C(I)



C(II)

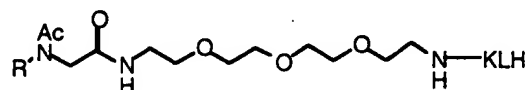


C(III)

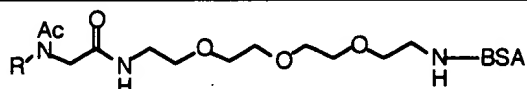
R—HN—KLH

Number	R
1	A(I)
2	B(I)
3	C(I)

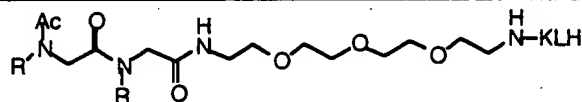
Number	
4	
5	



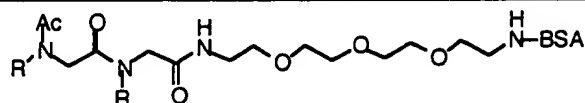
Number	R	Number	R	Number	R
6	A(I)	9	B(I)	12	C(I)
7	A(II)	10	B(II)	13	C(II)
8	A(III)	11	B(III)	14	C(III)



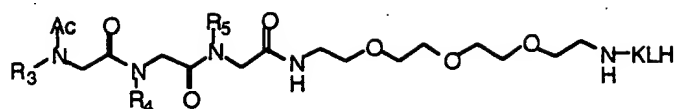
Number	R	Number	R	Number	R
15	A(I)	18	B(I)	21	C(I)
16	A(II)	19	B(II)	22	C(II)
17	A(III)	20	B(III)	23	C(III)



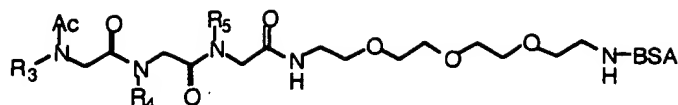
Number	R	Number	R	Number	R
15	A(I)	18	B(I)	21	C(I)
16	A(II)	19	B(II)	22	C(II)
17	A(III)	20	B(III)	23	C(III)



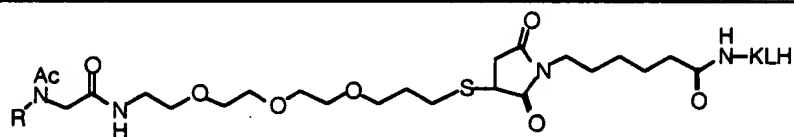
Number	R	Number	R	Number	R
24	A(I)	27	B(I)	30	C(I)
25	A(II)	28	B(II)	31	C(II)
26	A(III)	29	B(III)	32	C(III)



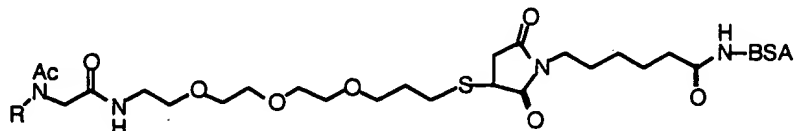
Number	R ₃	R ₄	R ₅	Number	R ₃	R ₄	R ₅
33	A(I)	A(I)	A(I)	41	C(III)	C(III)	C(III)
34	A(II)	A(II)	A(II)	42	A(II)	B(II)	A(II)
35	A(III)	A(III)	A(III)	43	B(II)	A(II)	B(II)
36	B(I)	B(I)	B(I)	44	A(II)	C(II)	A(II)
37	B(II)	B(II)	B(II)	45	C(II)	A(II)	C(II)
38	B(III)	B(III)	B(III)	46	B(II)	C(II)	B(II)
39	C(I)	C(I)	C(I)	47	C(II)	B(II)	C(II)
40	C(II)	C(II)	C(II)	48	A(II)	B(II)	C(II)



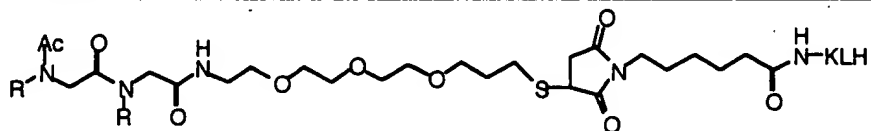
Number	R_3	R_4	R_5	Number	R_3	R_4	R_5
49	A(I)	A(I)	A(I)	57	C(III)	C(III)	C(III)
50	A(II)	A(II)	A(II)	58	A(II)	B(II)	A(II)
51	A(III)	A(III)	A(III)	59	B(II)	A(II)	B(II)
52	B(I)	B(I)	B(I)	60	A(II)	C(II)	A(II)
53	B(II)	B(II)	B(II)	61	C(II)	A(II)	C(II)
54	B(III)	B(III)	B(III)	62	B(II)	C(II)	B(II)
55	C(I)	C(I)	C(I)	63	C(II)	B(II)	C(II)
56	C(II)	C(II)	C(II)	64	A(II)	B(II)	C(II)



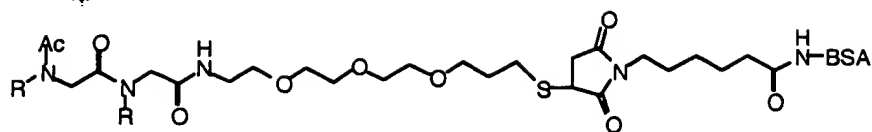
Number	R	Number	R	Number	R
65	A(I)	68	B(I)	71	C(I)
66	A(II)	69	B(II)	72	C(II)
67	A(III)	70	B(III)	73	C(III)



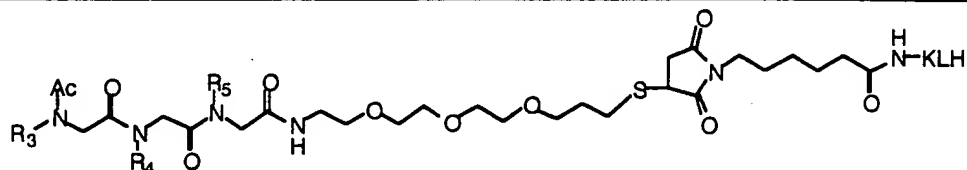
Number	R	Number	R	Number	R
74	A(I)	77	B(I)	80	C(I)
75	A(II)	78	B(II)	81	C(II)
76	A(III)	79	B(III)	82	C(III)



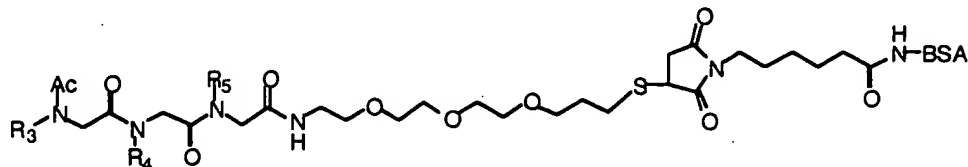
Number	R	Number	R	Number	R
83	A(I)	86	B(I)	89	C(I)
84	A(II)	87	B(II)	90	C(II)
85	A(III)	88	B(III)	91	C(III)



Number	R	Number	R	Number	R
92	A(I)	95	B(I)	98	C(I)
93	A(II)	96	B(II)	99	C(II)
94	A(III)	97	B(III)	100	C(III)

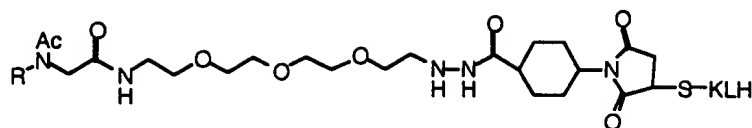


Number	R ₃	R ₄	R ₅	Number	R ₃	R ₄	R ₅
101	A(I)	A(I)	A(I)	109	C(III)	C(III)	C(III)
102	A(II)	A(II)	A(II)	110	A(II)	B(II)	A(II)
103	A(III)	A(III)	A(III)	111	B(II)	A(II)	B(II)
104	B(I)	B(I)	B(I)	112	A(II)	C(II)	A(II)
105	B(II)	B(II)	B(II)	113	C(II)	A(II)	C(II)
106	B(III)	B(III)	B(III)	114	B(II)	C(II)	B(II)
107	C(I)	C(I)	C(I)	115	C(II)	B(II)	C(II)
108	C(II)	C(II)	C(II)	116	A(II)	B(II)	C(II)



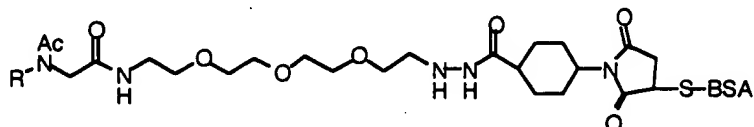
Number	R ₃	R ₄	R ₅	Number	R ₃	R ₄	R ₅
117	A(I)	A(I)	A(I)	125	C(III)	C(III)	C(III)
118	A(II)	A(II)	A(II)	126	A(II)	B(II)	A(II)
119	A(III)	A(III)	A(III)	127	B(II)	A(II)	B(II)
120	B(I)	B(I)	B(I)	128	A(II)	C(II)	A(II)
121	B(II)	B(II)	B(II)	129	C(II)	A(II)	C(II)
122	B(III)	B(III)	B(III)	130	B(II)	C(II)	B(II)
123	C(I)	C(I)	C(I)	131	C(II)	B(II)	C(II)
124	C(II)	C(II)	C(II)	132	A(II)	B(II)	C(II)

Table 15



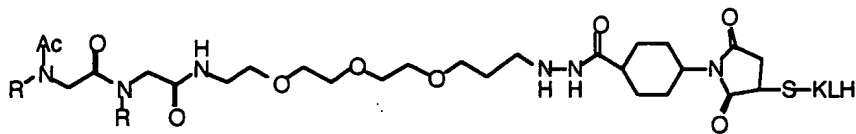
Number	R	Numbe	R	Numbe	R
133	A(I)	136	B(I)	139	C(I)
134	A(II)	137	B(II)	140	C(II)
135	A(III)	138	B(III)	141	C(III)

Table 16



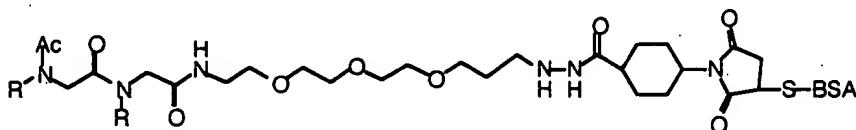
Numbe	R	Numbe	R	Numbe	R
142	A(I)	145	B(I)	148	C(I)
143	A(II)	146	B(II)	149	C(II)
144	A(III)	147	B(III)	150	C(III)

Table 17



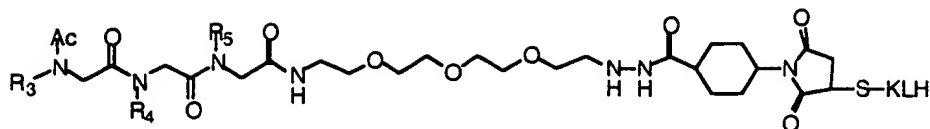
Numbe	R	Numbe	R	Numbe	R
151	A(I)	154	B(I)	157	C(I)
152	A(II)	155	B(II)	158	C(II)
153	A(III)	156	B(III)	159	C(III)

Table 18



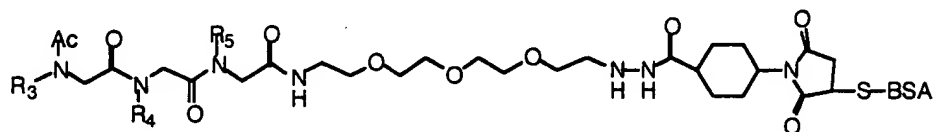
Numbe	R	Numbe	R	Numbe	R
160	A(I)	163	B(I)	166	C(I)
161	A(II)	164	B(II)	167	C(II)
162	A(III)	165	B(III)	168	C(III)

Table 19



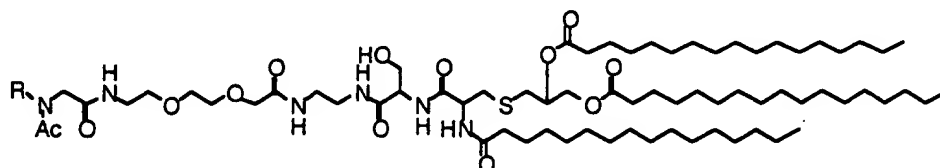
Numbe	R ₃	R ₄	R ₅	Numbe	R ₃	R ₄	R ₅
169	A(I)	A(I)	A(I)	177	C(III)	C(III)	C(III)
170	A(II)	A(II)	A(II)	178	A(II)	B(II)	A(II)
171	A(III)	A(III)	A(III)	179	B(II)	A(II)	B(II)
172	B(I)	B(I)	B(I)	180	A(II)	C(II)	A(II)
173	B(II)	B(II)	B(II)	181	C(II)	A(II)	C(II)
174	B(III)	B(III)	B(III)	182	B(II)	C(II)	B(II)
175	C(I)	C(I)	C(I)	183	C(II)	B(II)	C(II)
176	C(II)	C(II)	C(II)	184	A(II)	B(II)	C(II)

Table 20



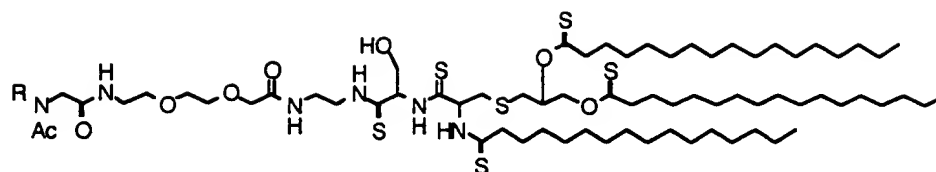
Numbe	R ₃	R ₄	R ₅	Numbe	R ₃	R ₄	R ₅
185	A(I)	A(I)	A(I)	193	C(III)	C(III)	C(III)
186	A(II)	A(II)	A(II)	194	A(II)	B(II)	A(II)
187	A(III)	A(III)	A(III)	195	B(II)	A(II)	B(II)
188	B(I)	B(I)	B(I)	196	A(II)	C(II)	A(II)
189	B(II)	B(II)	B(II)	197	C(II)	A(II)	C(II)
190	B(III)	B(III)	B(III)	198	B(II)	C(II)	B(II)
191	C(I)	C(I)	C(I)	199	C(II)	B(II)	C(II)
192	C(II)	C(II)	C(II)	200	A(II)	B(II)	C(II)

Table 21



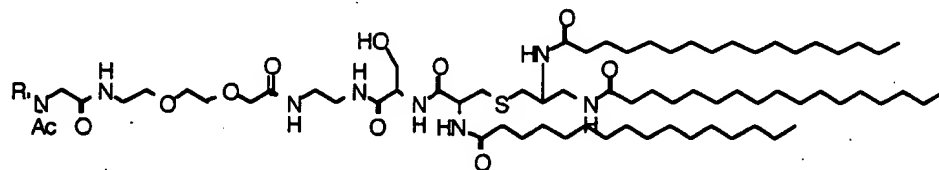
Numbe	R	Numbe	R	Numbe	R
201	A(I)	204	B(I)	207	C(I)
202	A(II)	205	B(II)	208	C(II)
203	A(III)	206	B(III)	209	C(III)

Table 22



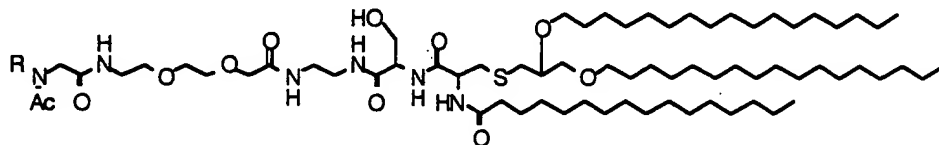
Numbe	R	Numbe	R	Numbe	R
210	A(I)	213	B(I)	216	C(I)
211	A(II)	214	B(II)	217	C(II)
212	A(III)	215	B(III)	218	C(III)

Table 23



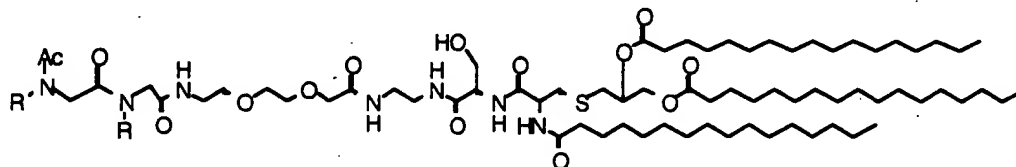
Numbe	R	Numbe	R	Numbe	R
219	A(I)	222	B(I)	225	C(I)
220	A(II)	223	B(II)	226	C(II)
221	A(III)	224	B(III)	227	C(III)

Table 24



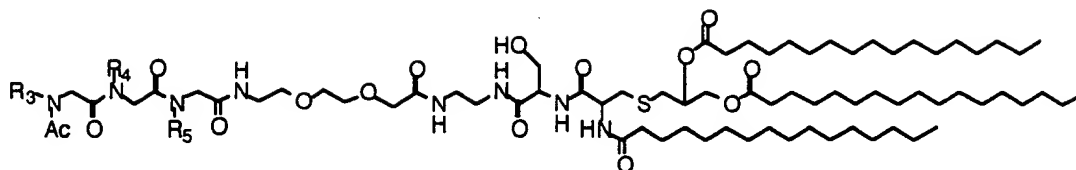
Numbe	R	Numbe	R	Numbe	R
228	A(I)	231	B(I)	234	C(I)
229	A(II)	232	B(II)	235	C(II)
230	A(III)	233	B(III)	236	C(III)

Table 25



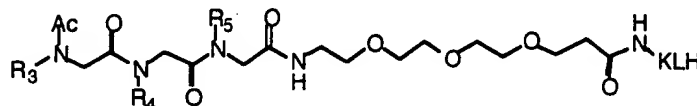
Numbe	R	Numbe	R	Numbe	R
237	A(I)	240	B(I)	243	C(I)
238	A(II)	241	B(II)	244	C(II)
239	A(III)	242	B(III)	245	C(III)

Table 26



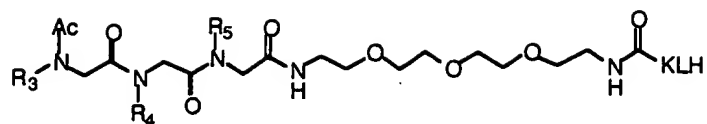
Numbe	R ₃	R ₄	R ₅	Numbe	R ₃	R ₄	R ₅
246	A(I)	A(I)	A(I)	254	C(III)	C(III)	C(III)
247	A(II)	A(II)	A(II)	255	A(II)	B(II)	A(II)
248	A(III)	A(III)	A(III)	256	B(II)	A(II)	B(II)
249	B(I)	B(I)	B(I)	257	A(II)	C(II)	A(II)
250	B(II)	B(II)	B(II)	258	C(II)	A(II)	C(II)
251	B(III)	B(III)	B(III)	259	B(II)	C(II)	B(II)
252	C(I)	C(I)	C(I)	260	C(II)	B(II)	C(II)
253	C(II)	C(II)	C(II)	261	A(II)	B(II)	C(II)

Table 27



Numbe	R ₃	R ₄	R ₅	Numbe	R ₃	R ₄	R ₅
262	A(I)	A(I)	A(I)	270	C(III)	C(III)	C(III)
263	A(II)	A(II)	A(II)	271	A(II)	B(II)	A(II)
264	A(III)	A(III)	A(III)	272	B(II)	A(II)	B(II)
265	B(I)	B(I)	B(I)	273	A(II)	C(II)	A(II)
266	B(II)	B(II)	B(II)	274	C(II)	A(II)	C(II)
267	B(III)	B(III)	B(III)	275	B(II)	C(II)	B(II)
268	C(I)	C(I)	C(I)	276	C(II)	B(II)	C(II)
269	C(II)	C(II)	C(II)	277	A(II)	B(II)	C(II)

Table 28



Numbe	R ₃	R ₄	R ₅	Numbe	R ₃	R ₄	R ₅
278	A(I)	A(I)	A(I)	286	C(III)	C(III)	C(III)
279	A(II)	A(II)	A(II)	287	A(II)	B(II)	A(II)
280	A(III)	A(III)	A(III)	288	B(II)	A(II)	B(II)
281	B(I)	B(I)	B(I)	289	A(II)	C(II)	A(II)
282	B(II)	B(II)	B(II)	290	C(II)	A(II)	C(II)
283	B(III)	B(III)	B(III)	291	B(II)	C(II)	B(II)
284	C(I)	C(I)	C(I)	292	C(II)	B(II)	C(II)
285	C(II)	C(II)	C(II)	293	A(II)	B(II)	C(II)

PHARMACOLOGICAL EXPERIMENT

Immunization and antiserum preparation

Vaccine used to immunize were prepared as below. Glycoprotein antigen (ex. 1mg) suspended in phosphate buffered saline (ex. 1mg) were mixed with equivalent volume of adjuvant (ex. Freund complete adjuvant, and BCG etc.). Female BALB/c mice (6 weeks of age) were subcutaneously immunized with 200 μ l/mouse of vaccine. Mice were injected on days 0, 14, 28, and bled 1 week after the 3rd immunization. Antiserum (-) was obtained from the blood centrifuged at 1,200 \times g for 20min.

Measurement of antibody titer.

Microtiter 96-well plate were coated with Tn antigen. IgG and IgM antibody titers were measured by ELISA with horse anti-mouse IgG antibody and anti-mouse IgM antibody, respectively, as second antibody. Human colon carcinoma cell line LS-174T cells cultured in microtiter 96-well plates, and were immobilized with methanol. As described, IgG and IgM antibody titers were measured by ELISA. Effects of each compounds described below on antibody titer were evaluated by this assay.

1		3	
4		5	
6		12	
15		33 (a)	
33 (b)		33 (c)	

IgG and IgM antibody titer (against Tn antigen) in mouse serum after vaccination were shown in table 30 and 31 respectively. IgG and IgM antibody titer (against LS-174T cell) after vaccination were shown in table 32.

Table 30

			1st immunization	2nd immunization	3rd immunization
Number	dose	Adjuvant	IgG	IgG	IgG
1	10 μ g	BCG	Not tested	Not Tested	2667 \pm 3233
3	10 μ g	BCG	800 \pm 0	2160 \pm 1431	4960 \pm 4879
5	10 μ g	BCG	Not tested	Not tested	2640 \pm 1252
6	10 μ g	BCG	200 \pm 200	1600 \pm 980	7680 \pm 10207
12	1 μ g	BCG	600 \pm 283	1920 \pm 1213	7680 \pm 10207
15	10 μ g	BCG	560 \pm 358	2080 \pm 1073	6400 \pm 5879
33(a)	1 μ g	BCG	200 \pm 346	1760 \pm 1315	6880 \pm 10516
33(b)	10 μ g	BCG	240 \pm 220	1920 \pm 720	7700 \pm 3320
33(c)	10 μ g	BCG	1040 \pm 780	1760 \pm 1320	4840 \pm 5010

Table 31

			1st immunization	2nd immunization	3rd immunization
Number	dose	Adjuvant	IgM	IgM	IgM
1	10 μ g	BCG	Not tested	Not tested	1333 \pm 462
3	10 μ g	BCG	40 \pm 89	1280 \pm 1213	2920 \pm 3257
5	10 μ g	BCG	Not tested	Not tested	1040 \pm 537
6	10 μ g	BCG	0 \pm 0	360 \pm 297	600 \pm 616
12	1 μ g	BCG	160 \pm 358	2880 \pm 716	1600 \pm 980
15	10 μ g	BCG	100 \pm 173	960 \pm 358	5140 \pm 2817
33	1 μ g	BCG	0 \pm 0	660 \pm 313	1860 \pm 2617
33(b)	10 μ g	BCG	100 \pm 100	1280 \pm 440	1960 \pm 2530
33(c)	10 μ g	BCG	10400 \pm 5400	3400 \pm 130	400 \pm 250

Table 32

Number	dose	Adjuvant	IgG	IgM
3	10 μ g	BCG	2400 \pm 1131	1520 \pm 1073
5	10 μ g	BCG	800 \pm 0	2160 \pm 1431
5	10 μ g	BCG	640 \pm 590	1120 \pm 438
6	10 μ g	BCG	1120 \pm 1242	1200 \pm 560
33(a)	1 μ g	BCG	1520 \pm 1073	1440 \pm 358

Antibody depended cell mediated cytotoxic response (ADCC)

LS-174T cells used as target cells and mononuclear cells from peripheral blood in human were used as effector cells. The target cells were seeded into microtiter 96-well plate (1×10^3 cells/ well/ 50 μ l),

and were added $0.5 \mu\text{Ci}$ / well of $^{51}\text{CrCl}_2$. the cells supernatants were harvested and counted in a gamma counter. The cytotoxicity was calculated as the percentage of releasable counts subtracting the spontaneous release. The results were shown in Table 33.

Table 33

Dilution	cpm/1000cells
200	913
400	685
800	318
1600	281
3200	103
6400	46

Purified carrier protein

Keyhole limpet hemocyanin (KLH, CHEMICONINTERNATIONAL INC.) was purified by the previously published method. KLH (500mg) was suspended in 50ml phosphate buffered saline (PBS(-)) and centrifuged at $1,200 \times g$ for 20 min. Resulted supernatant was centrifuged at $43,000 \times g$ for 15min. Resulted sediment was suspended in PBS(-) and further centrifuged at $43,000 \times g$ for 15min and resulted sediment was used as carrier protein.

Immunization

C-linked Tn-KLH conjugate or C-linked sTn-KLH conjugate (1 to $10 \mu\text{g}$) were immunized subcutaneously to female BALB/c mice with BCG($50 \mu\text{g}$) 3 times at two weeks interval. One week after the last immunization, mice were anesthetized and bloods were collected from abdominal vein. Antisera were separated by centrifugation, and IgG or IgM antibody titers against gp120 were assayed by ELISA. The titer was defined as the highest dilution yielding an absorbance of greater over that of normal sera. The results were shown in Table 34.

Table 34

Number	IgG	IgM
3	12,800	6,400
5	12,800	12,800
1 2	12,800	6,400
1 5	6,400	12,800
3 3	6,400	6,400

Our results also shows the potent immunogenicity of metabolic and catabolic stable "C-glycopeptoid" with or even without carrier protein. On other hand, Danishefsky's team reported the O-Tn, O-STn, O-TF antigens dose not have less potent immunogenicity themselves, but attached to carrier proteins such as KLH. (S.J.Danishefsky et al, 1998, 120, 1427-14285.)

We firstly showed the concept and efficacy of using "C-glycopeptoid" for the promising immunotherapy of cancer and HIV.

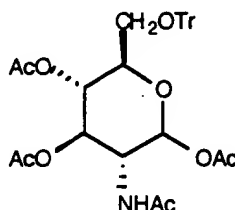
EXAMPLE

The following Examples are provide only for the purpose of the preparation of the compounds and not restrict to the disclosed invention.

Referencial Example 1

The preparation of 2-Acetylamino-1,3,4-tri-O-acetyl-6-O-triphenylmethyl-2-deoxy- α -D-glucopyranose

(compound 1a-2)



A suspension of N-acetylglucosamine (200g, 0.9mol) and trityl chloride (250g, 0.9mol) in pyridine (363ml) was heated to 85°C. After the suspension was dissolved, acetic anhydride (280ml, 2.97mol) was added and stirred for 23h at room temperature. The reaction mixture was slowly poured into ice water - acetic acid. The mixture was stirred for 3h and the resulting precipitate was collected, followed by washing with water. 400g (75%) of the objective compound was obtained.

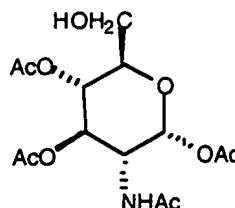
MS (m/z) : 590, 531, 452, 243, 165.

IR (cm⁻¹) neat : 3364, 1749, 1656, 1218.

¹H-NMR (CDCl₃) δ : 3.02 (1H, dd, $J=10.8, 3.9\text{Hz}$) , 3.27 (1H, dd, $J=10.3, 2.0\text{Hz}$) , 3.87 (1H, ddd, $J=9.3, 2.0, 2.0\text{Hz}$) , 4.54 (1H, ddd, $J=11.2, 9.3, 3.9\text{Hz}$) , 5.17 (1H, dd, $J=11.2, 9.8\text{Hz}$) , 5.35 (1H, dd, $J=9.8, 9.8\text{Hz}$) , 5.53 (1H, d, $J=8.8\text{Hz}$) , 6.29 ((1H, d, $J=3.4\text{Hz}$), 7.31-7.17 (9H, m) , 7.41-7.43 (6H, m) .

Referencial Example 2

The preparation of 2-acetylamino-1,3,4-tri-O-acetyl-2-deoxy- α -D-glucopyranose (compound 1a-3)



The trityl compound (168g) obtained from the above mentioned Referencial Example 1 was dissolved

in diethylether (420ml), then formic acid (420ml) was added at room temperature and the mixture was stirred for 7h. After the reaction was finished, the reaction mixture was poured into ice cold water and neutralized by NaHCO₃, followed by removal of diethylether, and the resulting precipitate was filtrated. The filtrate was extracted with chloroform. After drying (Na₂SO₄), the solvent was removed under reduced pressure. 46g (46%) of the objective alcohol was obtained.

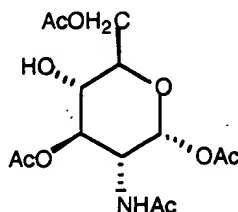
MS (m/z) : 347, 304, 228, 114.

IR (cm⁻¹) neat : 3280, 3076, 1749, 1665, 1221.

¹H-NMR (CDCl₃) δ : 1.91 (3H, s) , 2.04 (3H, s) , 2.16 (3H, s) , 3.55 (1H, dd, J=12.8, 4.4Hz) , 3.66 (1H, dd, J=12.8, 2.2Hz) , 3.78 (1H, ddd, J=10.1, 4.3, 2.2Hz) , 4.43 (1H, ddd, J=10.9, 9.0, 3.6Hz) , 5.14 (1H, t, J=9.7Hz) , 5.25 (1H, dd, J=10.8, 9.6Hz) , 5.76 (1H, d, J=9.0Hz) , 6.15 (1H, d, J=3.6Hz) .

Referencial Example 3

The preparation of 2-acetyl-amino-1,3,6-tri-O-acetyl-2-deoxy-α-D-glucopyranose (compound 1a-4)



To a solution of the primary alcohol compound (81g, 0.23mol) obtained from the above mentioned Referencial Example 2 in toluene (1600ml) was added acetic acid (16ml) and the mixture was stirred at 80°C for 15h. After the reaction was completed, the reaction mixture was concentrated under reduced pressure. The redsidue was purified by silicagel column chromatography (AcOEt). 99g (58%) of the objective compound was obtained as a colorless oil.

MS (m/z) : 347, 304, 262, 228, 114.

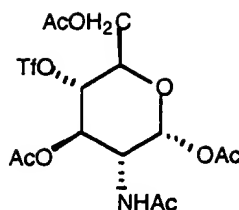
IR (cm⁻¹) neat : 3370, 3010, 1737, 1659, 1230.

¹H-NMR (CDCl₃) δ : 1.93 (3H, s) , 2.13 (3H, s) , 2.17 (3H, s) , 3.70 (1H, dd, J=9.8, 3.5Hz) , 3.84 (2H, d, J=3.5Hz) , 3.90 (1H, dd, J=9.8, 9.2Hz) , 4.30 (1H, ddd, J=11.1, 9.0, 3.7Hz) , 5.12 (1H, dd, J=11.1, 9.2Hz) , 5.71 (1H, d, J=9.0Hz) , 6.11 (1H, d, J=3.6Hz) .

Referencial Example 4

The preparation of 2-acetyl-amino-1,3,6-tri-O-acetyl-4-trifluoromethanesulfonyl-2-deoxy-α-D-

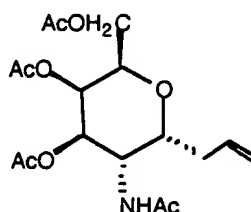
glucopyranose
(compound 1a-5)



The alcohol compound (5.0g, 14.3mmol) obtained from the above mentioned Referencial Example 3 was dissolved in dichloromethane (50ml) and added pyridine (5ml). The solution was cooled to -40°C , then triflic anhydride (3.1ml 18.7mmol) was added dropwise to the mixture. After stirring for 2h, the reaction mixture was poured into ice cold water and extracted with dichloromethane. The organic layer was washed with 10% HCl and dried (Na_2SO_4). The solvent was removed under reduced pressure. 7.83g of the objective compound was obtained as a colorless oil.

Referencial Example 5

The preparation of 3-(2-acetyl-3,4,6-tetra-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-propene (compound 1a-6)



To 2.0g (9.0mmol) of N-acetylgalactosamine was slowly added acetyl chloride (4.0ml) at 0°C . The mixture was stirred for 14h at room temperature. After the reaction, the mixture was poured into ice cold water and extracted with chloroform. The organic layer was neutralized by satd. NaHCO_3 , and washed with water and brine. After drying (Na_2SO_4), the solvent was removed under reduced pressure. 3.3g of N-acetylamino-1-Chloro-tri-O-acetyl-2-deoxy-galactosamine was obtained as a colorless oil. To a solution of the obtained compound (3.3g, 9.0mmol) in toluene, was added allyltributyltin (8.5ml) and 2, 2'-azobisisobutyronitrile (AIBN) (0.25g) under argon atmosphere. The reaction mixture was heated to 80°C and stirred for 6h. After the reaction was completed, the mixture was cooled to room temperature. The solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt: n-hexane=4:1). 0.85g (25.4%) of the oily objective compound was obtained as a colorless oil.

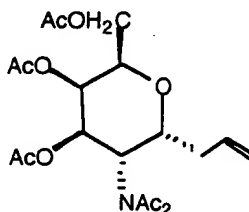
Mass (m/e) : 371, 330, 210, 150, 101, 59.

IR (cm⁻¹) KBr : 3290, 3071, 1746, 1658, 1020.

¹H-NMR(C₆D₆) δ : 1.47 (3H, s), 1.63 (3H, s), 1.66 (3H, s), 1.67 (3H, s), 1.99 (1H, m), 2.19 (1H, m), 3.94 (1H, m), 4.26 (1H, d, d, J=3, 5Hz), 4.37 (2H, m), 4.83 (1H, m), 5.00 (2H, m), 5.17 (1H, d, J=7Hz), 5.43 (1H, t, J=3Hz), 5.68 (1H, m), 6.19 (1H, S).

Referencial Example 6

The preparation of 3-(2-diacetyl-amino-3,4,6-tetra-O-acetyl-2-deoxy-α-D-galactopyranosyl)-1-propene
(compound 1a-9)



To a solution of the compound (1.5g, 4.0mmol) obtained from the above mentioned Referencial Example 5 in isopropenyl acetate (15ml) was added *p*-toluensulfonic acid (20mg). The reaction mixture was stirred at 55°C for 42h. After the mixture was cooled to room temperature, triethylamine was added and stirred for 15min. The mixture was concentrated. The residue was purified by silicagel column chromatography (BW-200, AcOEt: n-hexane=1:1). 1.0g (66%) of the objective diacetate compound was obtained as a colorless oil.

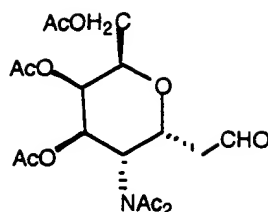
Mass (m/e) : 413, 372, 330, 270, 210, 179, 150, 126, 101, 59.

IR (cm⁻¹) KBr : 3050, 1749, 1668, 1233, 780

¹H-NMR(CDCl₃) δ : 1.95 (3H, s), 2.03 (3H, S), 2.16 (3H, s), 2.17 (1H, m), 2.39 (3H, s), 2.75 (1H, m), 4.05 (2H, m), 4.15 (2H, m), 4.61 (1H, d, d, J=4,8Hz), 5.11 (2H, m), 5.50 (1H, d, J=3Hz), 5.75 (1H, m), 5.95 (1H, dd, J=3,11Hz).

Referencial Example 7

The preparation of 3-(2-acetyl-amino-3,4,6-tetra-O-acetyl-2-deoxy-α-D-galactopyranosyl)-1-acetaldehyde
(compound 1a-10)



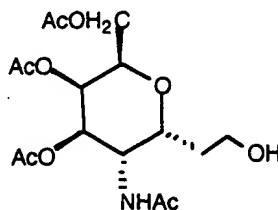
To a solution of the compound (0.74g, 1.78mmol) obtained from the above mentioned Referencial Example 6 in tetrahydrofrane (10ml) was added water (10ml), NaIO₄ (1.9g, 8.91mmol) and 4% OsO₄ solution under an atmosphere of argon. The mixture was stirred for 4h at room temperature. After the reaction was completed, the reaction mixture was extracted with ethyl acetate and washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. 0.77 g (98%) of the objective aldehyde compound was obtained as a colorless oil.

IR (cm⁻¹) KBr : 1746, 1371, 1230, 1054, 665.

¹H-NMR(CDCl₃) δ : 1.95 (3H, s), 2.04 (3H, s), 2.16 (3H, s), 2.37 (6H, s), 2.85 (1H, m), 3.17 (1H, dd, J=2,8Hz), 4.11 (3H, m), 4.75 (2H, m), 5.54 (1H, m), 5.81 (1H, d,d, J=3.5,11Hz), 9.67 (1H,s).

Referencial Example 8

The preparation of 3-(2-acetyl-amino-3,4,6-tetra-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-ethanol (compound 1a-11)



To a solution of the compound (0.77g, 1.85mmol) obtained from the above mentioned Referencial Example 7 in methanol (10ml) was added sodium borohydride (0.1g, 2.78mmol) at 0°C and the mixture was stirred for 10min. The reaction mixture was poured into satd.NH₄Cl and extracted with dichloromethane. The organic layer was washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt: MeOH=10:1). 0.25g (36%) of the oily objective compound was obtained as a colorless oil.

Mass (m/e) : 357(M⁺), 316, 238, 183, 141, 101, 59.

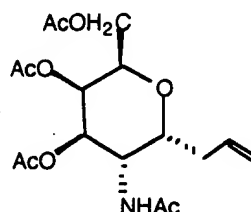
IR (cm⁻¹) KBr : 1743, 1680, 1398, 1236.

¹H-NMR(CDCl₃) δ : 1.60 (1H, m), 1.95 (1H, m), 2.00 (3H, s), 2.09 (3H, m), 2.10 (3H, s), 2.12

(3H, s), 3.17 (1H, dd, $J=3$, 8Hz) 3.76 (2H, m), 4.05-4.18 (3H, m), 4.42 (3H, m), 5.32 (1H, t, $J=3$ Hz), 5.73 (1H, d, $J=8$ Hz).

Referencial Example 9

The preparation of 3-(2-acetyl-amino-3,4,6-tetra-O-acetyl-2-deoxy- α -D-glucopyranosyl)-1-propene (compound 1b-2)



To N-acetylglucosamine 100g (0.45mol) was added acetyl chloride (200ml) at 0°C and stirred for 23h. After the reaction, the mixture was extracted with chloroform and the mixture was poured into ice cold water and stirred for 10min. The organic layer was neutralized by satd. NaHCO₃ and dried (Na₂SO₄). The solvent was removed under reduced pressure. Diethyl ether was added to the residue and the resulting precipitate was collected. 117g (71%) of 2-acetyl-amino-1-chloro-3,4,6-tetra-O-acetyl-2-deoxy- α -D-glucose was obtained as a colorless solid. To a solution of the obtained compound (78g, 0.21mol) in tetrahydrofuran (400ml) was added allyltributyltin (198ml, 0.64mol) and 2,2'-azobisisobutyronitrile (AIBN) (3.4g, 0.02mol). The reaction mixture was heated to 80°C and stirred for 16h under argon atmosphere. The reaction mixture was concentrated under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt: n-hexane=4:1). The mixture of allyl compound (1.62g) was obtained. To a solution of the obtained mixture in aceton (10ml) was added 1% HCl (6ml) and stirred for 2h. The mixture was concentrated under reduced pressure and the residue was extracted with chloroform (30ml). The organic layer was neutralized by satd. NaHCO₃ and dried (Na₂SO₄). The solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt: n-hexane=4:1). 73g (92%) of the objective compound was obtained as a colorless solid.

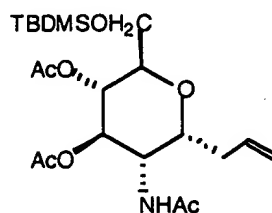
MS (m/z) : 371, 330, 312, 210, 126.

IR (cm^{-1}) neat : 3290, 3071, 1746, 1658, 1020.

¹H-NMR (CDCl₃) δ : 1.47 (3H, s) , 1.63 (3H, s) , 1.66 (3H, s) , 1.67 (3H, s) , 1.99 (1H, m) , 2.19 (1H, m) , 3.94 (1H, m) , 4.26 (1H, dd, $J=3$, 5Hz) , 4.37 (2H, m) , 4.38 (1H, m) , 5.01 (2H, m) , 5.17 (1H, d, $J=7$ Hz) , 5.43 (1H, t, $J=3$ Hz) , 5.68 (1H, m) , 6.19 (1H, s) .

Referencial Example 10

The preparation of 3-(2-acetylamino-3,4-di-O-acetyl-2-deoxy- α -D-glucopyranosyl)-1-propene (compound 1b-4)



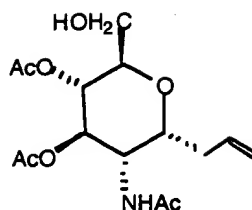
To a solution of the acetate compound (73g, 0.2mol) obtained from the above mentioned Referencial Example 9 in methanol (400ml) was added sodium methoxide (5g, 0.95mmol) at 0°C and stirred for 90min. After the reaction was completed, the reaction mixture was neutralized by IR-120 resin, filtrated and concentrated. 54.8g of the triol compound was obtained as a colorless solid. To a solution of the obtained triol compound (54.8g, 224mmol) in N,N-dimethylformamide (224ml) was added imidazole (30.8g, 448mmol), tert-butyldimethylsilyl chloride (40.5g, 268mmol) and dimethylaminopyridine (2.7g, 22.4mmol) and the mixture was stirred for 70h at 35°C. The reaction mixture poured into water and extracted with chloroform. The organic layer was neutralized by satd. NaHCO₃ and dried (Na₂SO₄). The solvent was removed under reduced pressure and 120g of the silyl compound was obtained. To the obtained silyl compound was added pyridine (108ml, 1.34mol), acetic anhydride (84.7ml, 0.89mol) and dimethylaminopyridine (13.7g, 0.11mol). The reaction mixture was stirred for 1h. After the reaction was finished, the mixture was poured into water and extracted with ethyl acetate. After drying (Na₂SO₄), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt: n-hexane=2:1). 33.4g (35%) of the objective alcohol compound was obtained as a colorless oil.

MS (m/z) : 428, 386, 326, 117.

¹H-NMR (CDCl₃) δ : 0.84 (9H, s) , 1.92 (3H, s) , 2.00 (3H, s) , 2.02 (3H, s) , 2.03 (3H, s) , 2.18-2.25 (1H, m) , 2.33-2.39 (1H, m) , 3.69 (2H, s) , 4.04-4.20 (3H, m) , 4.93-5.11 (4H, m) , 5.71-5.86 (2H, m) .

Referencial Example 11

The preparation of 3-(2-acetylamino-3,4-di-O-acetyl-2-deoxy- α -D-glucopyranosyl)-1-propene (compound 1b-6)



A solution of the silyl compound (10g, 23.1mmol) obtained from the above mentioned Referencial Example 10 in a mixture of tetrahydrofrane (10ml), acetic acid (30ml) and water (10ml) was stirred for 62h at 30°C. The reaction mixture was poured into water and extracted with chloroform, the organic extract was neutralized by satd.NaHCO₃, and dried over Na₂SO₄. The solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt). 7.5g (100%) of the objective alcohol compound was obtained as a colorless solid.

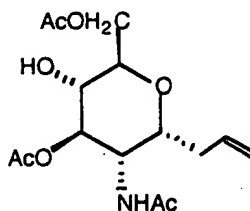
MS (m/z) : 330, 288, 268, 228, 126, 101.

IR (cm⁻¹) KBr : 3352, 2926, 1734, 1656, 1233.

¹H-NMR (CDCl₃) δ : 1.96 (3H, s) , 2.05 (3H, s) , 2.08 (3H, s) , 2.28-2.35 (1H, m) , 2.43-2.49 (1H, m) , 3.57-3.69 (3H, m) , 4.26-4.32 (2H, m) , 4.97 (1H, dd, J=8.3, 8.3Hz) , 5.10-5.19 (3H, m) , 5.78-5.86 (2H, m) .

Referencial Example 12

The preparation of 3-(2-acetylamino-3,6-di-O-acetyl-2-deoxy-α-D-glucopyranosyl)-1-propene (compound 1b-7)



A mixture of the primary alcohol compound (7.5g, 23.1mmol) obtained from the above mentioned Referencial Example 11 and acetic acid (0.75ml) in toluene (75ml) was stirred for 18h at 80°C. The mixture was concentrated under reduced pressure and the resulting residue was purified by silicagel column chromatography (AcOEt). 5.24g (70%) of the objective compound was obtained as a colorless oil.

MS (m/z) : 330, 228, 209, 168, 126, 101, 83.

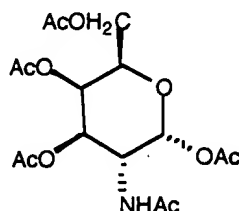
IR (cm⁻¹) KBr : 3352, 1734, 1656, 1233.

¹H-NMR (CDCl₃) δ : 1.95 (3H, s) , 2.13 (3H, s) , 2.14 (3H, s) , 2.30-2.36 (1H, m) ,

2.40-2.49 (1H, m) 、 3.55-3.59 (1H, m) 、 3.66-3.70 (1H, m) 、 4.18 (1H, dd, $J=12.2$, 2.9Hz) 、 4.22-4.29 (2H, m) 、 4.51 (1H, dd, $J=12.2$, 4.9Hz) 、 4.99 (1H, dd, $J=8.3$, 9.7Hz) 、 5.10-5.16 (2H, m) 、 5.72-5.82 (1H, m) 、 5.90 (1H, d, $J=8.3$ Hz) .

Example 1

The preparation of 2-acetylamino-1,3,4,6-tetra-O-acetyl-2-deoxy- α -D-galactopyranose (compound 1a-7)



To a solution of cesium acetate (13.7g, 71.5mmol) in dimethylsulfoxide (15ml) was added a solution of the triflate compound (7.83g) obtained from the above mentioned Referencial Example 4 in dimethylsulfoxide (15ml). After the mixture was stirred for 3h, the mixture was concentrated under reduced pressure. The residue was poured into water and extracted with dichloromethane, then dried over Na_2SO_4 . The solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt). 3.4g (61%) of the objective compound was obtained as a colorless oil.

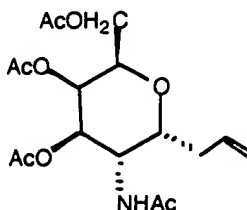
MS (m/z) : 389、330、287、241、114.

IR (cm^{-1}) neat : 1746、1656、1218、1128.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.88 (3H, s) 、 1.96 (3H, s) 、 1.97 (3H, s) 、 2.10 (3H, s) 、 3.99 (1H, dd, $J=11.2$, 6.6Hz) 、 4.04 (1H, dd, $J=11.2$, 6.8Hz) 、 4.20 (1H, ddd, $J=6.8$, 6.6, 0.9Hz) 、 4.63 (1H, ddd, $J=11.6$, 9.0, 3.6Hz) 、 5.14 (1H, dd, $J=11.7$, 3.2Hz) 、 5.36 (1H, dd, $J=3.1$, 0.7Hz) 、 5.82 (1H, d, $J=9.0$ Hz) 、 6.15 (1H, d, $J=3.6$ Hz) .

Example 2

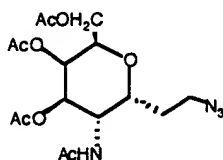
The preparation of 3-(2-acetylamino-3,4,6-tetra-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-propene (compound 1a-8)



The alcohol compound (13.2g, 40.1mmol) obtained from the above mentioned Referencial Example 12 was dissolved in a mixture of dichloromethane (130ml) and pyridine(13ml). Then triflic anhydride (8.1ml, 48.1mmol) was added dropwise at -40°C and stirred for 4h. The mixture was poured into ice cold water and extracted with dichloromethane and the organic extract was washed with 10%HCl and dried over Na_2SO_4 . The solvent was removed under reduced pressure and 16.1g of triflate compound (16.1g) was obtained. A solution of the obtained triflate compound (16.1g) in dimethylsulfoxide (60ml) was added to a solution of cesium acetate (20.0g, 104mmol) in dimethylsulfoxide (100ml). After the mixture was stirred for 3h, the mixture was concentrated under reduced pressure. The residue was poured into water and extracted with dichloromethane, then dried over Na_2SO_4 . The solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt). 10.9g (84%) of the objective compound was obtained as a colorless solid.

Example 3

The preparation of 2-(2-acetylamino-3,4,6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-ethylazide (compound 2-1a)



To a solution of the alcohol compound (2.33g, 6.22mmol) obtained from the above mentioned Referencial Example 8 in tetrahydrofuran (62ml) was added diphenylphosphoryl azide (2.68ml, 12.4mmol) and triphenylphosphine (3.25g, 12.4mmol). The solution was cooled to 0°C , diisopropyl azodicarboxylate (2.44ml, 12.4mmol) was added slowly to the solution and the mixture was stirred for 1h. The reaction mixture was concentrated under reduced pressure and the resulting residue was purified by silicagel column chromatography (AcOEt:benzene=1:1). 1.92g (77%) of the objective compound was obtained as a colorless oil.

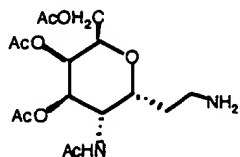
MS (m/e) : 401, 357, 313, 277, 166, 101.

IR (cm^{-1}) neat : 3244, 3046, 2092, 1737, 1656.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.66-1.72 (1H, m) , 1.83-1.89 (1H, m) , 2.00 (3H, s) , 2.07 (3H, s) , 2.08 (3H, s) , 2.12 (3H, s) , 3.35-3.39 (2H, m) , 4.02-4.12 (2H, m) , 4.31-4.35 (2H, m) , 4.45 (1H, ddd, $J=8.3, 8.3, 4.9\text{Hz}$) , 5.14 (1H, dd, $J=8.8, 3.4\text{Hz}$) , 5.33 (1H, dd, $J=3.4, 3.4\text{Hz}$) , 6.23 (1H, d, $J=8.3\text{Hz}$) .

Example 4

The preparation of 2-(2-acetylamino-3,4,6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-ethylamine (compound 2-2a)



The azide compound (982mg, 2.46mmol) obtained from the above mentioned Example 3 was dissolved in methanol (10ml), acetic acid (0.1ml) and 10% Pd-C (98mg) were added to the solution. The reaction mixture was stirred for 88h under an atmosphere of H₂. The suspension was filtered and the filtrate was concentrated under reduced pressure. The resulting residue was purified by silicagel column chromatography (CHCl₃:MeOH:H₂O=8:2:0.2). 662mg (72%) of the objective compound was obtained as a colorless oil.

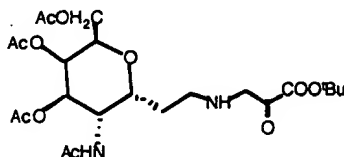
MS (m/e) : 374, 317, 256, 166, 115.

IR (cm⁻¹) neat : 3280, 2932, 1740, 1656.

¹H-NMR (CD₃OD) δ : 1.75-1.79 (1H, m) , 1.97-2.01 (1H, m) , 1.99 (3H, s) , 2.03 (3H, s) , 2.05 (3H, s) , 2.09 (3H, s) , 3.02-3.04 (2H, m) , 4.07 (1H, dd, $J=11.7$, 4.4Hz) , 4.18-4.19 (1H, m) , 4.31-4.45 (3H, m) , 5.12 (1H, dd, $J=9.3$, 3.4Hz) , 5.42 (1H, dd, $J=3.4$, 3.4Hz) .

Example 5

The preparation of t-butyl-2-({2-[2-acetylamino-3,4,6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl]ethyl}amino) acetate (compound 2-3a)



To a solution of the amine compound (590mg, 1.58mmol) obtained from the above mentioned Example 4 in dichloromethane (15.8ml) was added triethylamine (0.33ml, 2.73mmol) and tert-butyl bromoacetic acid (0.35ml, 2.37mmol). After the mixture was stirred for 2h at 60°C, the mixture was concentrated under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt:MeOH = 10:1). 225mg (27%) of the objective amide compound was obtained as a colorless oil.

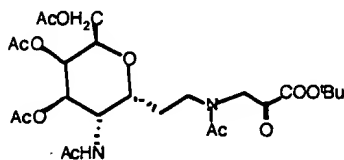
MS (m/e) : 489, 414, 387, 224, 164, 88.

IR (cm⁻¹) neat : 3328, 1740, 1656, 1233.

¹H-NMR (CD₃OD) δ : 1.54 (9H, s) , 1.61-1.65 (1H, m) , 1.96-1.98 (1H, m) , 1.96 (3H, s) , 2.02 (3H, s) , 2.03 (3H, s) , 2.10 (3H, s) , 2.62-2.77 (2H, m) , 3.28-3.37 (2H, m) , 4.10 (1H, dd, J=10.7, 4.9Hz) , 4.16 (1H, ddd, J=8.3, 8.3, 2.4Hz) , 4.22 (1H, ddd, J=8.3, 8.3, 3.4Hz) , 4.24-4.32 (1H, m) , 4.40 (1H, dd, J=9.8, 4.9Hz) , 5.12 (1H, dd, J=9.8, 2.9Hz) , 5.40 (1H, dd, J=2.9, 2.9Hz) .

Example 6

The preparation of t-butyl 2-(N-{2-[2-acetylamino-3,4,6-tri-O-acetyl-2-deoxy-α-D-galactopyranosyl] ethyl} acetamino) acetate (compound 2-4a)



The amine compound (100mg, 0.205mmol) obtained from the above mentioned Example 5 was dissolved in pyridine (1ml), acetic anhydride (0.039ml, 0.41mmol) and dimethylaminopyridine (12mg, 0.103mmol) were added to the solution. After the solution was stirred for 1h, the mixture was poured into water and extracted with ethyl acetate, the organic layer was washed with satd. CuSO₄ and brine, and dried over Na₂SO₄. The solvent was removed under reduced pressure, then the resulting residue was purified by silicagel column chromatography (AcOEt:MeOH=20:1). 100mg (92%) of the objective compound was obtained as a colorless oil.

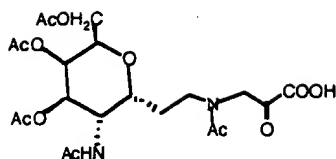
MS (m/e) : 530, 487, 429, 387, 222, 57.

IR (cm⁻¹) neat : 2968, 1740, 1650, 1230.

¹H-NMR (CD₃OD) δ : 1.45 (9H, s) , 1.73-1.77 (1H, m) , 1.92-1.97 (1H, m) , 1.97 (3H, s) , 2.00 (3H, s) , 2.03 (3H, s) , 2.10 (3H, s) , 2.16 (3H, s) , 3.40-3.60 (2H, m) , 3.89-4.30 (6H, m) , 4.40-4.44 (1H, m) , 5.07-5.14 (1H, m) , 5.38-5.40 (1H, m).

Example 7

The preparation of 2-(N-{2-[2-acetylamino-3,4,6-tri-O-acetyl-2-deoxy-α-D-galactopyranosyl] ethyl} acetamino) acetic acid (compound 2-5a)



A mixture of the ester compound (90mg, 0.17mmol) obtained from the above mentioned Example 6 and trifluoroacetic acid (0.2ml) in dichloromethane (1ml) was stirred for 3h. The mixture was concentrated under reduced pressure and the resulting residue was purified by silicagel column chromatography (CHCl₃:MeOH:AcOH=18:2:1). 70mg (87%) of the objective compound was obtained as a colorless oil.

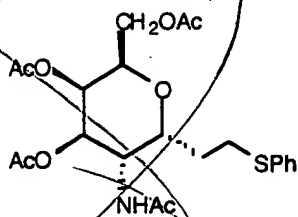
MS (m/e) : 474, 429, 314, 222, 69.

IR (cm⁻¹) neat : 1740, 1370, 1230.

¹H-NMR (CD₃OD) δ : 1.76-1.82 (1H, m) , 1.92-1.97 (1H, m) , 1.99 (3H, s) , 2.03 (3H, s) , 2.11 (3H, s) , 2.14 (3H, s) , 2.17 (3H, s) , 3.70-3.52 (2H, m) , 4.00-4.30 (6H, m) , 4.43-4.46 (1H, m) , 5.08-5.14 (1H, m) , 5.38-5.40 (1H, m) , 4.43-4.46 (1H, m) , 5.08-5.14 (1H, m) , 5.38-5.40 (1H, m).

Example 8

The preparation of 3-(2-acetyl-amino-3,4-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-phenylthioethane (compound 4-1)



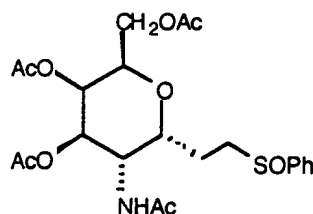
The compound (0.25g, 0.67mmol) obtained from the above mentioned Example 7 was dissolved in pyridine (3ml), tributylphosphine (0.42ml) and diphenyldisulfide (0.32g) were added to the solution. The mixture was stirred for 3h at 60°C under argon atmosphere. The reaction mixture was extracted with ethyl acetate and washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt:n-Hexane=10:1). 0.18g (56%) of the objective thiophenyl compound was obtained as a colorless oil.

Mass (m/e) : 467 (M⁺).

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.63 (1H, m), 1.94 (3H, s), 1.96 (1H, m), 2.03 (3H, s), 2.06 (3H, s), 2.56 (3H, s), 2.91 (1H, m), 3.24 (1H, m), 3.98 (1H, m), 4.51 (2H, m), 4.32 (1H, m), 4.42 (2H, m), 5.07 (1H, dd, $J=4, 9\text{Hz}$), 5.29 (1H, t, $J=3\text{Hz}$), 5.55 (1H, d $J=7\text{Hz}$), 7.21-7.38 (5H, m).

Example 9

The preparation of 3-(2-acetylamino-3,4-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-phenylsulfonyl-ethane (compound 4-2)



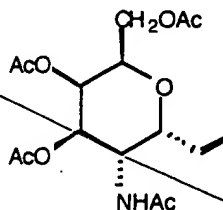
To a solution of the compound (0.14g, 0.29mmol) obtained from the above mentioned Example 8 in dichloromethane (2ml) was slowly added a solution of 3-chloroperoxybenzoic acid in dichloromethane (1.0ml) at -78°C . After stirring for 30min, diethylether (10ml) and 10% NaOH (1ml) was added to the reaction mixture and the mixture was stirred for 15min. The organic layer was separated and washed with water and brine. After drying (MgSO_4), the solvent was removed under reduced pressure. 0.15g (99%) of the objective compound was obtained as a colorless oil.

Mass (m/e) : 483 (M^+).

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.89 (1H, m), 1.91 (3H, s), 1.95 (3H, s), 2.05 (3H, s), 2.09 (1H, m), 1.96 (1H, m), 2.58 (1H, m), 2.80 (1H, t, $J=8\text{Hz}$), 3.01 (1H, m), 3.80 (1H, m), 3.95-4.10 (2H, m), 4.20 (1H, m), 4.35 (1H, m), 4.56 (2H, m), 5.10 (1H, dd, $J=4, 9\text{Hz}$), 5.27 (1H, t, $J=3\text{Hz}$), 6.50 (1H, d, $J=8\text{Hz}$), 7.4-7.60 (5H, m).

Example 10

The preparation of 3-(2-acetylamino-3,4-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-vinylene (compound 4-3)



A mixture of the compound (0.14g, 0.29mmol) obtained from the above mentioned Example 9 and

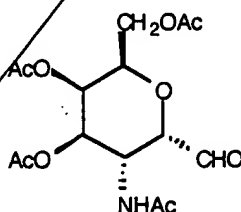
Cont
Al
diisopropylethylamine (0.09ml) in toluene (2ml) was refluxed for 18h. After the reaction mixture was cooled to room temperature, the mixture was extracted with ethyl acetate and washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt). 0.07g (70%) of the oily objective compound was obtained as a colorless oil.

Mass (m/e) : 357 (M⁺), 298, 255, 165, 101(BP), 59.

¹H-NMR(CDCl₃) δ : 1.96 (3H, s), 2.05 (3H, s), 2.06 (3H, s), 2.16 (3H, s), 4.14 (3H, m), 4.62 (1H, m), 4.76 (1H, m), 5.03 (1H, dd, J=4, 10Hz), 5.35 (1H, d, J=2Hz), 5.45 (3H, m), 5.95 (1H, m).

Example 11

The preparation of 3-(2-acetylamino-3,4-tri-O-acetyl-2-deoxy-α-D-galactopyranosyl)-1-carbaldehyde (compound 4-4)



Alm
As
To a mixture of the compound (0.07g, 0.20mmol) obtained from the above mentioned Example 10, in tetrahydrofran (2ml) and water was added NaIO₄ (0.16g, 0.78mmol) and 4% OsO₄ solution (0.01ml). After the mixture was stirred for 4h, the reaction mixture was extracted with ethyl acetate and washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. 0.705g (69.6%) of the objective aldehyde compound was obtained as a colorless oil.

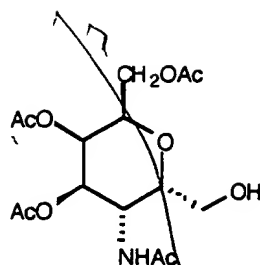
Mass (m/e) : 360(M⁺), 330, 300, 199, 131, 97(BP), 59.

¹H-NMR(CDCl₃) δ : 1.98 (3H, s), 2.06 (3H, s), 2.02 (3H, S), 2.17 (3H, s), 3.92 (1H, t, J=7Hz), 4.20 (2H, m), 4.59 (1H, d, J=Hz), 4.80 (1H, m), 5.09 (1H, dd, J=3, 9Hz), 5.38 (1H, d, J=3Hz), 6.22 (1H, d, J=9Hz), 9.83 (1H, S).

Example 12

Alm
As
The preparation of 3-(2-acetylamino-3,4-tri-O-acetyl-2-deoxy-α-D-galactopyranosyl)-1-methanol (compound 4-5)

Grub
a.s.



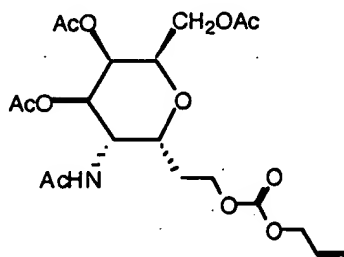
A mixture of the compound (0.77g, 1.85mmol) obtained from the above mentioned Example 11 and sodium borohydride (0.1g, 2.78mmol) in methanol (10ml) was stirred for 10min at 0°C. The reaction mixture was poured into satd. NH₄Cl and the mixture was extracted with dichloromethane, the organic layer was washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt). 0.25g (36%) of the objective alcohol compound was obtained as a colorless oil.

Mass (m/e) : 362(M⁺), 330, 300, 199, 139, 97(BP), 59.

¹H-NMR(CDCl₃) δ : 1.98 (3H, s), 2.06 (3H, s), 2.02 (3H, s), 2.17 (3H, s), 3.92 (1H, t, J=7Hz), 4.20 (2H, m), 4.59 (1H, d, J=3Hz), 4.80 (1H, m), 5.09 (1H, dd, J=3, 9Hz), 5.38 (1H, d, J=3Hz), 6.22 (1H, d, J=9Hz), 9.83 (1H, s).

Example 13

The preparation of 3-(2-acetamino-3,4,6-tetra-O-acetyl-2-deoxy-α-D-galactopyranosyl)-1-ethylvinylxyformate (compound 5-1a)



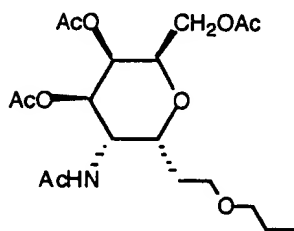
The compound (0.09g, 2.27mmol) obtained from the above mentioned Example 8 was dissolved in tetrahydrofuran (5ml), allyl chloroformate (0.026ml, 2.5mmol) was added to the solution in the presence of pyridine (1ml). After the solution was stirred for 30min, the reaction mixture was extracted with ethyl acetate and the organic layer was washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure and the resulting residue was purified by silicagel column chromatography (BW-200, AcOEt:n-hexane=4:1). 0.08g (70%) of the objective compound was obtained as a colorless oil.

IR (cm⁻¹) KBr : 1743, 1392, 1245, 1020.

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.58 (1H, m), 1.99, (3H, s), 2.07 (3H, s), 2.08 (3H, s), 2.12 (3H, s), 4.12 (2H, m), 4.19-4.35 (4H, m), 4.38 (1H, m), 4.48 (1H, m), 4.62 (2H, d, $J=6\text{Hz}$), 5.13 (1H, dd, $J=3, 8\text{Hz}$), 5.27-5.39 (3H, m), 5.64 (1H, d, $J=8\text{Hz}$).

Example 14

The preparation of 3-(2-acetamino-3,4,6-tetra-O-acetyl-2-deoxy- α -D-glactopyranosyl)-1-ethoxyprop-2-ene (compound 5-2a)



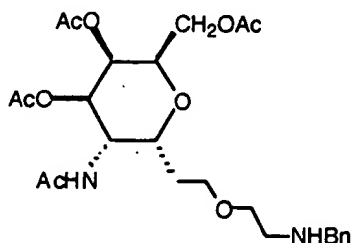
The compound (0.07g, 0.15mmol) obtained from the above mentioned Example 13 was dissolved in benzene (2ml), $\text{Pd}(\text{OAc})_2$ (0.7mg) and triphenylphosphine (4mg) were added to the solution under argon atmosphere. After the mixture was stirred for 2h at 70°C , the mixture was concentrated. The resulting residue was purified by silicagel column chromatography (BW-200, $\text{AcOEt}:\text{hexane}=4:1$). 0.045g (72.3%) of the objective compound was obtained as a colorless oil.

Mass (m/e) : 415(M^+), 358, 314, 277, 181, 152, 101, 59.

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.82 (1H, m), 1.91 (1H, m), 1.97 (3H, s), 2.05 (3H, s), 2.06 (3H, s), 2.12 (3H, s), 3.50 (2H, m), 3.97 (2H, d, $J=3\text{Hz}$), 4.06 (2H, m), 4.22 (1H, m), 4.28 (1H, m), 4.50 (1H, m), 5.12 (1H, dd, $J=3, 5\text{Hz}$), 5.15 (1H, dd, $J=2, 7\text{Hz}$), 5.30 (1H, m), 5.76 (1H, d, $J=8\text{Hz}$), 5.90 (1H, m).

Example 15

The preparation of 2-(2-acetyl-amino-3,4,6-tri O-acetyl-2-deoxy- α -D-glactopyranosyl)-1-[2-(benzylamino) ethoxy] ethane (compound 5-3a)



A solution of the compound (0.69g, 1.65mmol) obtained from the above mentioned Example 14 in methanol (5ml) and dichloromethane (5ml) was ozonized at -78°C. After the reaction was completed, dimethylsulfide was added to the mixture and the solution was stirred at room temperature. The mixture was concentrated and 0.69g (99%) of the aldehyde was obtained. To a solution of the obtained aldehyde in dichloromethane (5ml) was added benzylamine (0.22ml). After stirring for 15 min, sodium triacetoxyborohydride (0.5g) was added to the mixture and the mixture was stirred for 12h. The reaction mixture was extracted with chloroform and the organic layer was washed with water and brine. After drying (Na₂SO₄), the solvent was removed under reduced pressure and the resulting residue was purified by silicagel column chromatography (BW-200, chloroform:methanol=20:1). 0.51g (64.4%) of the objective compound was obtained as a colorless oil.

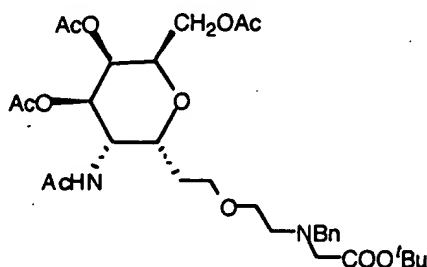
Mass (m/e) : 449 (M-NHAc), 383, 192, 120, 91.

IR (cm⁻¹) KBr : 3290, 2950, 1740, 1660, 1378, 1230, 1000.

¹H-NMR(CDCl₃) δ : 1.87 (1H, m), 1.95 (3H, s), 1.97 (1H, m), 2.05 (6H, m), 2.19 (3H, s), 2.81 (2H, t, J=5.0Hz), 3.51 (2H, m), 3.57 (2H, t, J=5.0Hz), 3.84 (1H, s), 4.03 (1H, m), 4.08 (1H, m), 4.38 (1H, m), 4.50 (1H, m), 5.13 (1H, dd, J=8.3, 2.1Hz), 5.31 (1H, t, J=3Hz), 5.85 (1H, d, J=8.0Hz), 7.32 (5H, m).

Example 16

The preparation of t-butyl 2-[(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl] ethoxy} ethyl) benzylamino] acetate (compound 5-4a)



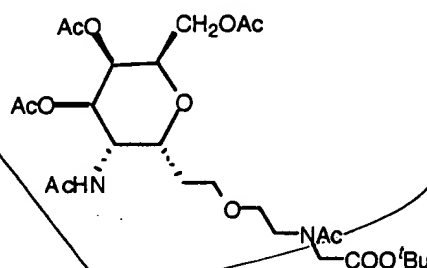
A mixture of the compound (0.51g, 1.03mmol) obtained from the above mentioned Example 15 and tert-butyl bromoacetic acid (0.3ml) in dichloromethane (5ml) was stirred for 16h at 60°C. After the reaction was completed, triethylamine was added to the mixture and stirred for 15min. The mixture was extracted with ethyl acetate and washed with water and brine. After drying (MgSO₄), the solvent was removed under reduced pressure and the resulting residue was purified by silicagel column chromatography (ABW-200, CHCl₃:MeOH=10:1). 0.23g (36.9%) of the objective compound was

obtained as a colorless oil.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.45 (9H, s), 1.75 (1H, m), 1.84 (1H, m), 1.97 (3H, s), 2.05 (3H, s), 2.06 (3H, s), 2.12 (3H, s), 2.86 (2H, t, $J=6.0\text{Hz}$), 3.29 (2H, s), 3.40-3.60 (4H, m), 3.83 (2H, s), 4.02 (1H, m), 4.11 (1H, m), 4.20 (1H, m), 4.32 (1H, m), 4.50 (1H, m), 5.11 (1H, dd, $J=6.0, 2.0\text{Hz}$), 5.31 (1H, t, $J=3\text{Hz}$), 5.70 (1H, d, $J=6.0\text{Hz}$), 7.33 (5H, m).

Example 17

The preparation of t-butyl 2-[(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethoxy} ethyl) amino] acetate (compound 5-5a)



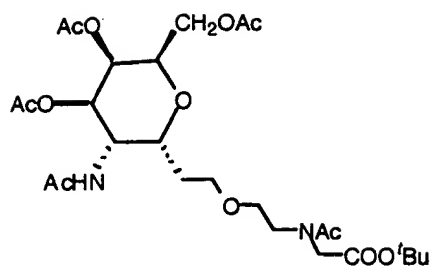
The compound (0.21g, 0.34mmol) obtained from the above mentioned Example 16 dissolved in methanol (10ml), acetic acid (0.5ml) and 10% Pd-C (20mg) were added to the solution. The reaction mixture was stirred for 3h under an atmosphere of H_2 , then the suspension was filtered through celite and the filtrate was concentrated. 0.18g (99%) of the objective compound was obtained as a colorless oil.

Mass (m/e) : 431, 373, 314, 91, 78.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.45 (9H, s), 1.91 (1H, m), 1.95 (1H, m), 2.01 (3H, s), 2.05 (6H, s), 2.05 (3H, s), 2.50 (1H, s), 2.87 (2H, t, $J=6.0\text{Hz}$), 3.40-3.70 (4H, m), 4.10 (2H, m), 4.20-4.41 (3H, m), 4.50 (2H, m), 5.20 (1H, dd, $J=6.0, 2.0\text{Hz}$), 5.33 (1H, t, $J=3\text{Hz}$), 6.05 (1H, d, $J=6.0\text{Hz}$).

Example 18

The preparation of t-butyl 2-[(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethoxy} ethyl) acetylamino] acetate (compound 5-6a)



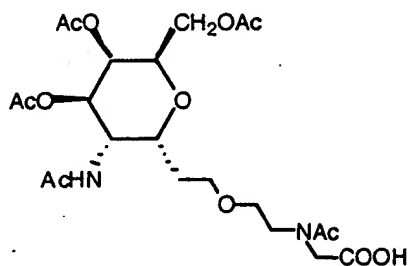
To a solution of the compound (0.18g, 0.34mmol) obtained from the above mentioned Example 17 in dichloromethane (5ml) was slowly added acetyl chloride (0.36ml) in the presence of diisopropylethylamine (0.1ml). After the solution was stirred for 2h, the mixture was concentrated under reduced pressure and the resulting residue was purified by silicagel column chromatography (BW-200, AcOEt). 0.13g (66.5%) of the objective compound was obtained as a colorless oil.

Mass (m/e) : 517 (M-Bu), 501, 431, 358, 314, 199, 144, 99, 72.

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.47 9H, s), 1.83 (1H, m), 1.93 (1H, m), 1.86 (3H, s), 1.91-2.20 (15H, m), 3.40-3.60 (10H, m), 3.96-4.40 (9H, m), 4.50 (1H, m), 5.18 (1H, dd, $J=6.0, 3.0$), 5.30 (1H, t, $J=3.0\text{Hz}$), 5.75 (1H, m).

Example 19

The preparation of 2-[N-(2-{2-[2-acetyl-amino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethoxy} ethyl) benzylamino] acetic acid (compound 5-7a)



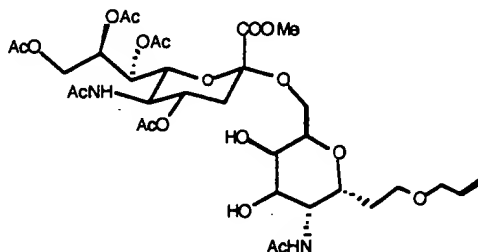
A mixture of the compound (0.15g, 0.26mmol) obtained from the above mentioned Example 18 and trifluoroacetic acid (0.4ml) in dichloromethane (2ml) was added to the mixture and stirred for 3h. The reaction mixture was concentrated and 0.13g (66.8%) of the objective compound was obtained as a colorless oil.

Mass (m/e) : 517 (M-Bu), 501, 431, 358, 314, 199, 144, 99, 72.

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 0.89 (1H, m), 0.97 (1H, m), 2.00 (3H, s), 2.03 (3H, s), 2.04 (3H, s), 2.15 (3H, s), 2.19 (3H, s), 3.40-3.65 (6H, m), 3.95-4.18 (3H, m), 4.30 (2H, m), 4.46 (1H, m), 5.10 (1H, d, $J=4.0\text{Hz}$), 5.18 (1H, m).

Example 20

The preparation of O-(methyl 5-acetyl-amino-4,8,9-tetra-O-acetyl-3,5-dideoxy- β -D-glycero-D-galacto-2-nuno pyranosynate)-(2 \rightarrow 6)-2-(2-acetyl-amino-3,4-di-O-acetyl-2-deoxy- α -D-galctopyranosyl)-1-(prop-2-enyloxy) ethane (compound 6-2a)



A mixture of the alcohol compound (173mg, 0.66mmol) obtained from the above mentioned Example 14 and MS4A in tetrahydrofuran (10ml) was added di-tert-butylpyridine (0.29ml) and AgOTf (337mg) and the mixture was stirred for 30min. After cooling to -78°C, a solution of the sialyl chloride (670mg, 0.66mmol) in tetrahydrofurane (8ml) was added dropwise to the mixture and the mixture was stirred for 28h. The suspension was filtered through Celite and the filtrate was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (CHCl₃:MeOH= 10:1). 81mg (18%) of the objective compound was obtained as a colorless oil.

MS (ESI, m/e) : 785 (M⁺) .

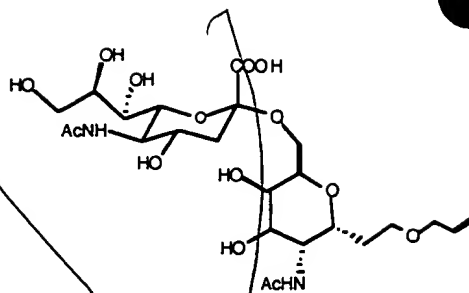
IR (cm⁻¹) : 3340, 2944, 1744, 1656.

¹H-NMR (CDCl₃) δ : 1.88 (3H, s) , 1.99 (3H, s) , 2.03 (3H, s) , 2.04 (3H, s) , 2.13 (3H, s) , 2.14 (3H, s) , 2.58 (1H, dd, J=17.4, 4.4Hz), 2.97 (1H, d, J=3.9Hz), 3.99-4.10 (4H, m) , 4.14-4.27 (2H, m) , 4.34 (1H, dd, J=12.2, 2.5Hz) , 4.40-4.43 (1H, m) , 4.84-4.91 (1H, m) , 5.21-5.39 (4H, m) , 5.87-5.96 (1H, m) , 6.74 (1H, d, J=5.9Hz) .

Example 21

The preparation of O-(methyl 5-acetyl-amino-3,5-dideoxy- β -D-glycero-D-galacto-2-nuno pyranosinate)-(2 \rightarrow 6)-2-(2-acetyl-amino-2-deoxy- α -D-galactopyranosyl)-1-(prop-2-enyloxy) ethane (compound 6-3a)

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Q13



A mixture of the compound (21mg, 0.027mmol) obtained from the above mentioned Example 20 and 2% K₂CO₃ (3ml) in methanol (9ml) was stirred for 20h. The reaction mixture was neutralized by 1% HCl, then the reaction mixture was concentrated under reduced pressure. The resulting residue was purified by silicagel column chromatography (PR-18, H₂O:AcOH=100:1). 13mg (81%) of the objective compound was obtained as a colorless oil.

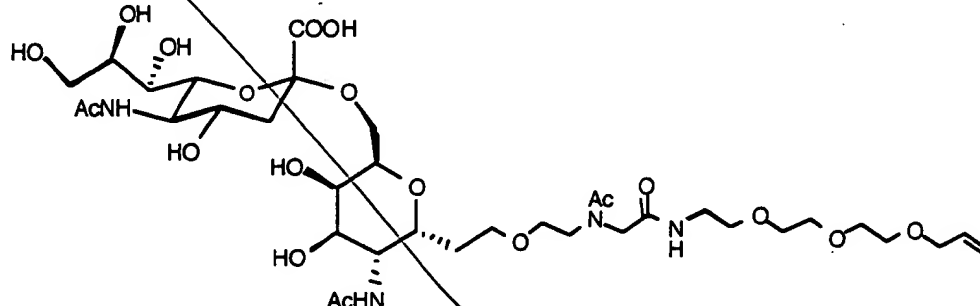
MS (ESI, m/e) : 579 (M-H)⁺

IR (cm⁻¹) neat : 3268, 1638, 1566.

¹H-NMR (CD₃OD) δ : 1.60-1.80 (2H, m) , 2.01-2.05 (1H, m) , 2.01 (3H, s) , 2.05 (3H, s) , 2.85-2.88 (1H, m) , 3.50-3.60 (3H, m) , 3.65-3.76 (5H, m) , 3.81-3.95 (5H, m) , 4.02 (2H, d, $J=5.4\text{Hz}$) , 4.21-4.33 (2H, m) , 5.18-5.21 (1H, m) , 5.29-5.34 (1H, m) , 5.91-6.00 (1H, m) .

Example 22

The preparation of O-(methyl 5-acetyl amino-3,5-dideoxy- β -D-glycero-D-galacto-2-nonuropysanosinate)-(2 \rightarrow 6)-2-[N-(2-{2-[2-acetyl amino-3,4,6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl]ethoxy}ethyl)acetyl amino]-N-(2-{2-[2-(2-oxoethoxy)ethoxy]ethoxy}ethyl)acetamide (6-3b)



To use of the compound obtained from the following mentioned Example 32, the objective compound was obtained according to the method described in Example 20-21.

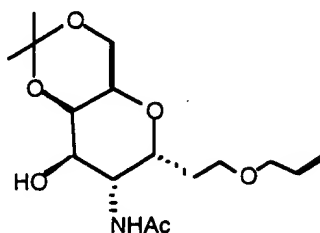
MS (ESI, m/e) : 877 (M+Na)⁺

IR (cm^{-1}) KBr : 3400, 2950, 1650, 1400, 1125.

$^1\text{H-NMR}$ (CD_3OD) δ : 1.62 (2H, m) , 2.01 (3H, s) , 2.04 (3H, s) , 2.20 (3H, s) , 2.81 (1H, dd, $J=2.8\text{Hz}$) , 3.40-3.98 (30H, m) , 4.18 (4H, m) , 4.21 (4H, m) , 5.20 (1H, d, $J=12\text{Hz}$) , 5.19 (1H, d, $J=12\text{Hz}$) , 6.98 (1H, m) .

Example 23

The preparation of the following compound.



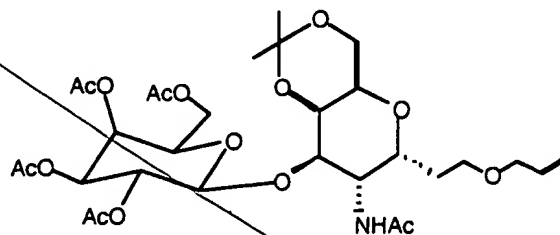
To a solution of the compound (0.12g, 0.41mmol) obtained from the above mentioned Example 14 in acetonitrile (10ml) was added benzaldehyddimethylacetal (0.12ml) and p-toluensulfonate (3.8mg) and the mixture was stirred for 6h at 60°C under argon atmosphere. After cooling to room temperature, the mixture was extracted with ethyl acetate and the organic layer was washed with water and brine. After drying (MgSO_4), the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt). 0.10g (64.4%) of the objective acetal compound was obtained as a colorless oil.

MS (ESI, m/e) : 377 (M^+)

$^1\text{H-NMR}$ (CDCl_3) δ : 1.96 (2H, m) , 1.99 (3H, s) , 3.43-3.60 (3H, m) , 3.75 (1H, d, $J=3\text{Hz}$) , 4.01 (3H, m) , 4.11 (2H, m) , 4.42 (2H, m) , 5.20 (2H, dd, $J=3, 8\text{Hz}$) , 5.60 (1H, s) , 5.90 (1H, m) , 6.20 (1H, d, $J=3\text{Hz}$) , 7.30-7.56 (5H, m)

Example 24

The preparation of the following compound.



A mixture of the compound (100mg, 0.41mmol) obtained from the above mentioned Example 23 and

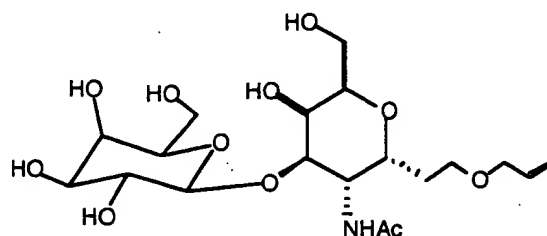
MS4A in dichloromethane (10ml) was added di tert-butylpyridine (0.12ml) and AgOTf (0.14g) and the mixture was stirred for 30min. After cooling to -78°C, a solution of the galactose derivatives (0.22g 0.41mmol) in dichloromethane was added dropwise to the mixture. After the reaction was completed, the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (BW-200, AcOEt). 0.10g (64.4%) of the objective compound was obtained as a colorless oil.

MS (ESI, m/e) : 707 (M)⁺

¹H-NMR (CDCl₃) δ : 1.96 (2H, m) , 1.99 (3H, s) , 3.43-3.60 (3H, m) , 3.75 (1H, d, J=3Hz) , 4.01 (3H, m) , 4.11 (2H, m) , 4.42 (2H, m) , 5.20 (2H, , dd, J=3, 8Hz) , 5.60 (1H, s) , 5.90 (1H, m) , 6.20 (1H, d, J=3Hz) , 7.30-7.56 (5H, m)

Example 25

The preparation of the following compound.



A solution of the compound (0.11g, 0.16mmol) obtained from the above mentioned Example 24 in 80% acetic acid was heated to 70°C and stirred for 2h. The solvent was removed under reduced pressure mixture and the obtained diol compound was dissolved in methanol (5ml). Sodium methoxide (2mg) was added to the solution and the mixture was stirred for 2h at room temperature. The reaction mixture was neutralized by Amberlite IR-120 and filtrated and the filtrate was concentrated under reduced pressure. 0.1g (64.4%) of the objective compound was obtained.

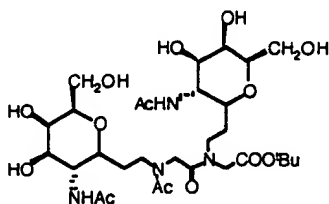
MS (ESI, m/e) : 451 (M)⁺

¹H-NMR (CDCl₃) δ : 1.75 (1H, m) , 2.00 (1H, m) , 2.13 (3H, s) , 3.16 (1H, m) , 3.55 (2H, m) , 3.60 (1H, m) , 3.69-3.82 (3H, m) , 3.98 (2H, m) , 4.01-4.10 (5H, m) , 4.30 (1H, m) , 4.80 (3H, m) , 5.11 (1H, m) , 5.18 (2H, m) , 5.23 (1H, m) , 5.40 (1H, m) , 5.90 (1H, m)

Example 26

The preparation of t-butyl 2-[N-(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-

glactopyranosyl] ethyl) -2-[N-(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethoxy} ethyl) acetylamino] acetate (compound 8-3a)



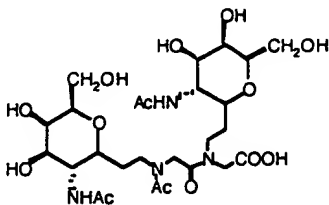
A mixture of carboxylic acid (67mg, 0.14mmol) and amine (69mg, 0.14mmol) obtained from above mentioned Example 7 and 5 were dissolved in acetonitrile (1.4ml), diisopropylethylamine (0.027ml) and O-(benzotriazol-1-yl) N, N, N', N'-tetramethylhydroniumtetrafluoroborate (TBTU) (50mg) were added to the mixture. After the reaction mixture was stirred for 24h, the mixture was poured into brine and extracted with chloroform, the organic layer was dried over Na₂SO₄ and the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography (AcOEt:MeOH=10:1). 72mg (54%) of the objective compound was obtained as a colorless oil.

MS (ESI, m/e) : 944 (M⁺) .

¹H-NMR (CD₃OD) δ : 1.46 (9H, s) , 1.77-1.83 (1H, m) , 1.92-1.97 (1H, m) , 1.90 (3H, s) , 1.91 (3H, s) , 1.94 (3H, s) , 1.95 (3H, s) , 2.00 (3H, s) , 2.01 (3H, s) , 2.04 (3H, s) , 2.05 (3H, s) , 2.11 (3H, s) , 3.41-3.68 (4H, m) , 3.90-4.59 (14H, m) , 5.09-5.16 (2H, m) , 5.40-5.42 (2H, m) .

Example 27

The preparation of 2-[N-(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl) -2-[N-(2-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethoxy} ethyl) acetylamino] acetic acid (compound 8-1b)



A solution of the ester compound (62mg, 65.7 μ mol) obtained from the above mentioned Example 26 and trifluoroacetic acid (0.2ml) in dichloromethane (1ml) was stirred for 4h. The reaction mixture was concentrated under reduced pressure, and the resulting residue was purified by silicagel column chromatography (CHCl₃:MeOH:AcOH=18:2:1). 50mg (86%) of the objective compound was

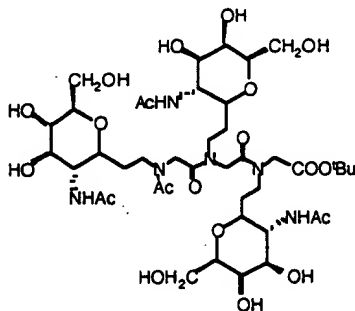
obtained as a colorless oil.

MS (ESI, m/e) : 888 (M^+) .

$^1\text{H-NMR}$ (CD_3OD) δ : 1.77-1.82 (2H, m) 、 1.94 (3H, s) 、 1.95 (3H, s) 、 1.97 (3H, s) 、 1.99 (3H, s) 、 2.00 (3H, s) 、 2.02 (3H, s) 、 2.04 (3H, s) 、 2.07 (3H, s) 、 2.11 (3H, s) 、 3.34-3.77 (4H, m) 、 4.06-4.87 ((14H, m) 、 5.10-5.15 (2H, m) 、 5.33-5.41 (2H, m) .

Example 28

The preparation of t-butyl N-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl} -2-(N-{2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl} acetylamino)-N-({N-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl}-N-[(N-carbamoyl)methyl]carbamoyl}methyl) acetate (compound 8-3b)



To a solution of carboxylic acid (48mg, 54.1 μ mol) and amine (26.3mg, 54.1 μ mol) obtained from above mentioned Example 27 and 17 in acetonitrile (1ml) was added diisopropylethylamine (10 μ l, 59.5 μ mol) and O-(benzotriazol-1-yl) N, N, N', N'-tetramethylhydroniumtetrafluoroborate (TBTU) (19mg, 59.5 μ mol). After stirring for 38h, the mixture was poured into brine and extracted with chloroform, the organic layer was dried over Na_2SO_4 and the solvent was removed under reduced pressure. The resulting residue was purified by silicagel column chromatography ($\text{AcOEt}:\text{MeOH}=5:1$). 40mg (54%) of the objective compound was obtained as a colorless oil.

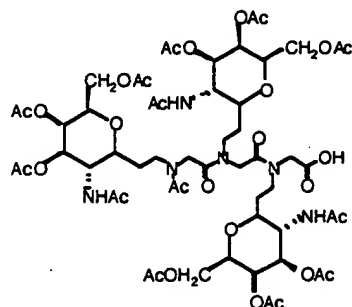
MS (ESI, m/e) : 1382 ($M+\text{Na}$) $^+$.

$^1\text{H-NMR}$ (CD_3OD) δ : 1.46 (9H, s) 、 1.73-1.83 (3H, m) 、 1.94-2.18 (42H, m) 、 3.41-3.78 (8H, m) 、 4.03-4.55 (21H, m) 、 5.10-5.15 (3H, m) 、 5.40-5.42 (3H, m) .

Example 29

The preparation of N-{2-[2-acetylamino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl} -

2-(N-{2-acetyl-amino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl} acetyl-amino)-N-({N-
 {2-[2-acetyl-amino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-glactopyranosyl] ethyl}-N-[(N-
 carbamoyl)methyl]carbamoyl}methyl) acetic acid (compound 8-1c)



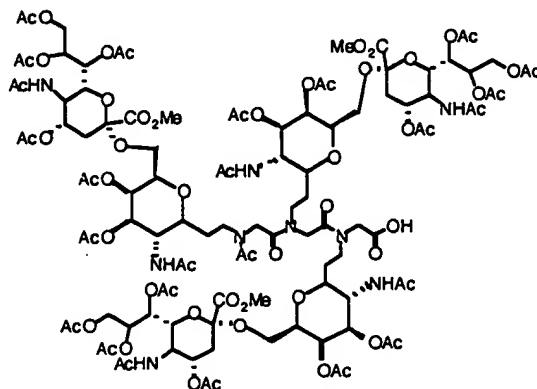
A solution of the ester compound (40mg, 29.5 μ mol) obtained from the above mentioned Example 28 and trifluoroacetic acid (0.2ml) in dichloromethane (1ml) was stirred for 16h. The reaction mixture was concentrated under reduced pressure and the resulting residue was purified by silicagel column chromatography (CHCl_3 :MeOH:AcOH=18:2:1). 18mg (47%) of the objective compound was obtained as a colorless oil.

MS (ESI, m/e) : 1302 (M^+) .

$^1\text{H-NMR}$ (CD_3OD) δ : 1.65-1.75 (3H, m) , 2.00-2.14 (42H, m) , 3.31-3.60 (6H, m) , 4.05-4.50 (21H, m) , 5.05-5.10 (3H, m) , 5.30-5.39 (3H, m) .

Example30

The preparation of the following compound.

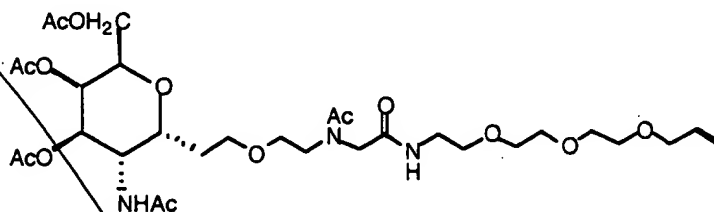


To use of the compound obtained from the above mentioned Example 20, the objective compound was obtained according to the method described in Example 26-28.

$^1\text{H-NMR}$ (CD_3OD) δ : 1.80-2.25 (69H, m), 3.01-3.68 (50H, m), 3.90-4.58 (6H, m).

Example 31

The preparation of 2-(2-Acetylamino-3,4,6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-(2-{N-[(N-{2-[2-(2-prop 2-enyloxyethoxy)ethoxy]ethyl}carbamoyl)methyl]acetylamino}ethoxy)ethane (compound 8-4)



To a solution of carboxylic acid (23mg, 44.4 μ mol) obtained from above mentioned Example 19 and amine (17mg, 88.8 μ mol) in acetonitrile (1ml) was added diisopropylethylamine (9 μ l, 48.8 μ mol), O-(benzotriazol-1-yl) N, N, N', N'-tetramethylhydroniumtetrafluoroborate (TBTU) (16mg, 48.8 μ mol). After the mixture was stirred for 4h, the mixture was poured into brine and extracted with chloroform, the organic layer was washed with 10% HCl and satd. NaHCO₃. After drying (Na₂SO₄), the solvent was removed under reduced pressure and the resulting residue was purified by silicagel column chromatography (AcOEt:MeOH=8:1). 20mg (65%) of the objective compound was obtained as a colorless oil.

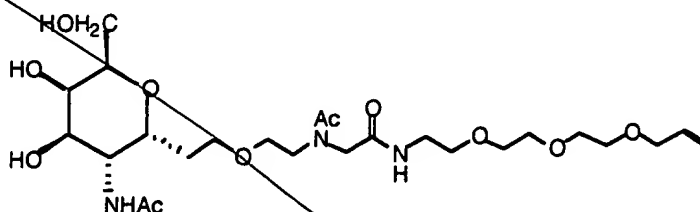
MS (ESI, m/e) : 688 (M^+).

IR (cm⁻¹) neat : 3286, 2860, 1743, 1650.

¹H-NMR (CDCl₃) δ : 1.80-1.86 (1H, m), 2.00 (3H, s), 2.06 (3H, s), 2.12 (3H, s), 2.13 (3H, s), 2.18 (3H, s), 3.42-3.67 (18H, m), 4.40-4.11 (6H, m), 4.20-4.30 (1H, m), 4.34-4.41 (1H, m), 4.42-4.47 (1H, m), 5.10-5.14 (1H, m), 5.17-5.20 (1H, m), 5.25-5.30 (1H, m), 5.31-5.32 (1H, m), 5.86-5.98 (1H, m).

Example 32

The preparation of 2-[N-(2-{2-[2-Acetylamino-2-deoxy- α -D-galactopyranosyl]ethoxy}ethyl)acetylamino]-N'-2-[2-(2-prop-2-oxyethoxy)ethoxy]ethyl]acetamide (compound 8-6)



A mixture of the acetate compound (19.5mg, 29.0 μ mol) obtained from the above mentioned Example 31 and sodium methoxide (3mg, 58.0 μ mol) in methanol (1ml) was stirred for 1.5h at 0°C. The reaction mixture was neutralized by IR-120, filtered and the filtrate was removed under reduced

pressure. 15.7mg (99%) of the objective triol compound was obtained as a colorless oil.

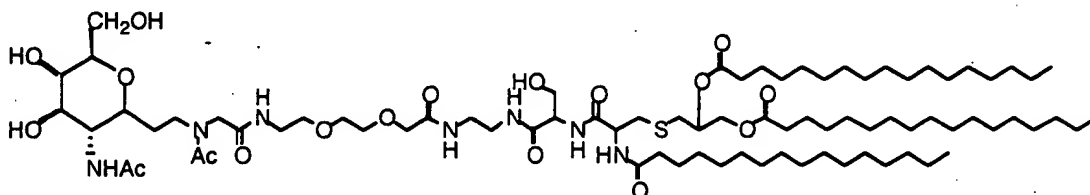
MS (ESI, m/e) : 562 (M^+).

IR (cm^{-1}) neat : 3272, 2932, 1636.

$^1\text{H-NMR}$ (CD_3OD) δ : 1.68-1.78 (1H, m) , 1.92-2.00 (1H, m) , 1.92 (3H, s) , 2.21 (3H, s) , 3.64-3.78 (24H, m) , 3.83-3.88 (1H, m) , 4.05-4.10 (2H, m) , 4.21-4.29 (2H, m) , 5.19-5.22 (1H, m) , 5.30-5.53 (1H, m) , 5.91-6.01 (1H, m) .

Example 33

The preparation of the following compound.



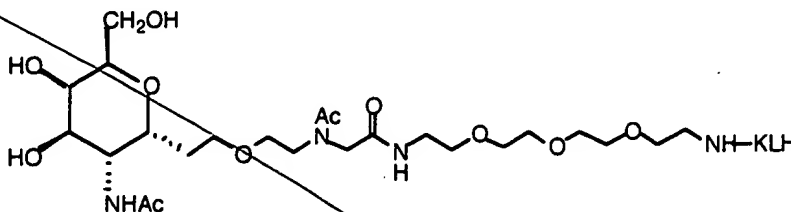
To a solution of the carboxylic acid (20mg, 57.5 μmol) obtained from the above mentioned Example 7 and amine (136mg, 115 μmol) in dimethylformamide (1ml) was added diisopropylethylamine (42 μl , 230 μmol), HATU (87mg, 230 μmol) and HOAt (16mg, 115 μmol). After the mixture was stirred for 24h, the mixture was poured into brine and extracted with chloroform, the organic layer was washed with 10% HCl and satd. NaHCO_3 . After drying (Na_2SO_4), the solvent was removed under reduced pressure and the resulting residue was purified by silicagel column chromatography ($\text{CHCl}_3:\text{MeOH}:\text{AcOH}=18:2:1$). 5mg (6%) of the objective compound was obtained as a colorless oil.

MS (ESI, m/e) : 1150

$^1\text{H-NMR}$ (CDCl_3 , ppm) : δ 1.70-2.12 (45H, m) , 3.41-3.79 (15H, m) , 4.09-4.58 (26H, m) , 5.08-5.36 (1H, m) , 5.40-5.48 (3H, m) , 5.90-6.05 (1H, m)

Example 34

The preparation of the following compound.

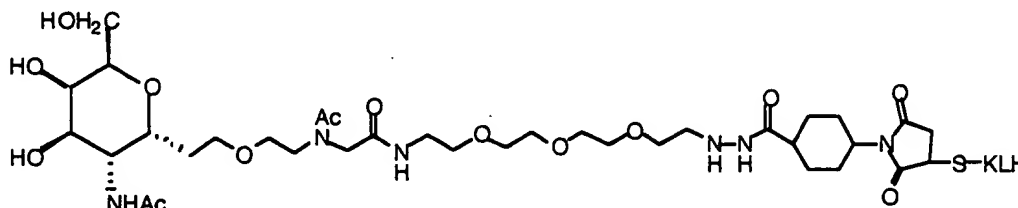


A solution of the compound obtained from the above in methanol and dichloromethane was ozonized at -78°C . The reaction mixture was treated with dimethylsulfide and concentrated to obtain the

aldehyde..To the mixture of this aldehyde and KLH in phosphate buffer was added sodium cyanoborohydride and stirred for 30h. After purified by dialysis using PBS(-), the objective glycoprotein antigen was obtained.

Example 35

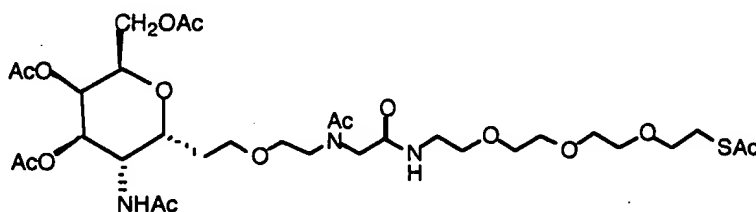
The preparation of the following compound.



The aldehyde obtained from the above mentioned Example 34 reacted with 4-(4-N-maleimidomethyl)cyclohexyl-1-carbonylhydrazine to obtain a maleimide derivative. To the mixture of this compound and KLH in phosphate buffer was added sodium cyanoborohydride. After purified by dialysis using PBS(-), the objective glycoprotein antigen was obtained.

Example 36

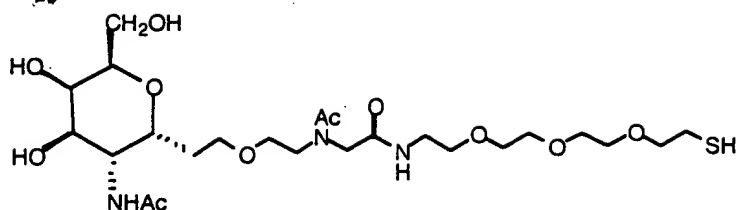
The preparation of 2-(2-acetyl-amino-3, 4, 6-tri-O-acetyl-2-deoxy- α -D-galactopyranosyl)-1-(2-{N-[(N-{2-[2-(3-acetylthiopropoxy)ethoxy]ethyl}carbonyl)methyl]acetyl-amino}ethoxy)ethane



To a solution of the olefine (22mg, 0.032mmol) obtained from the above mentioned Example 31 in dioxane (2ml) was added thioacetic acid (0.02ml) and the mixture was heated at 80°C for 6h. The solvent was removed under reduced pressure. The residue was purified by silicagel column chromatography (AcOEt:MeOH=9:1). 0.02g (82%) of the objective compound was obtained.

Example 37

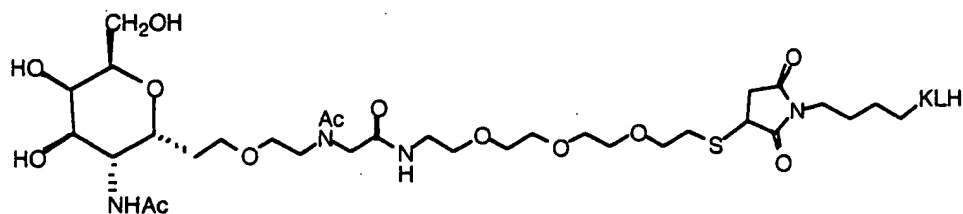
The preparation of 2-[N-(2-{2-[2-acetyl-amino-2-deoxy- α -D-galactopyranosyl] ethoxy} ethyl) acetyl-amino]-N-{2-[2-(2-sulfenylpropoxy)ethoxy]ethyl}acetamide



To a solution of the compound (20mg, 0.029mmol) obtained from the above mentioned Example 36 in methanol (1ml) was added sodium methoxide (2mg, 0.058mmol) and stirred for 12h. The reaction mixture neutralized by IR-120, filtered using celite, and the solvent was removed under reduced pressure. 8mg (51%) of the objective compound was obtained.

Example 38

The preparation of the following compound.



The compound obtained from the above mentioned Example 37 was added to maleimidated KLH with stirring and allowed to stand over 2h at 4°C. The reaction mixture was dialyzed with phosphate buffered saline (pH7.4) for 48h and with distilled water for 48h, followed by byophilization and obtained the objective compound.